Number 208



3. Quality Improvement
Interventions To Address
Health Disparities
Closing the Quality Gap:
Revisiting the State of the
Science

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# 3. Quality Improvement Interventions To Address Health Disparities

Closing the Quality Gap: Revisiting the State of the Science

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## **Preface**

The Agency for Healthcare Research and Quality (AHRQ), through its Evidence-based Practice Centers (EPCs), sponsors the development of evidence reports and technology assessments to assist public- and private-sector organizations in their efforts to improve the quality of health care in the United States. The reports and assessments provide organizations with comprehensive, science-based information on common, costly medical conditions, and new health care technologies and strategies. The EPCs systematically review the relevant scientific literature on topics assigned to them by AHRQ and conduct additional analyses when appropriate prior to developing their reports and assessments.

In 2004, AHRQ launched a collection of evidence reports, Closing the Quality Gap: A Critical Analysis of Quality Improvement Strategies, to bring data to bear on quality improvement opportunities. These reports summarized the evidence on quality improvement strategies related to chronic conditions, practice areas, and cross-cutting priorities.

This evidence report is part of a new series, Closing the Quality Gap: Revisiting the State of the Science. This series broadens the scope of settings, interventions, and clinical conditions, while continuing the focus on improving the quality of health care through critical assessment of relevant evidence. Targeting multiple audiences and uses, this series assembles evidence about strategies aimed at closing the "quality gap," the difference between what is expected to work well for patients based on known evidence and what actually happens in day-to-day clinical practice across populations of patients. All readers of these reports may expect a deeper understanding of the nature and extent of selected high-priority quality gaps, as well as the systemic changes and scientific advances necessary to close them.

AHRQ expects that the EPC evidence reports will inform consumers, health plans, other purchasers, providers, and policymakers, as well as the health care system as a whole, by providing important information to help improve health care quality.

We welcome comments on this evidence report or the series as a whole. Comments may be sent by mail to Shilpa Amin, M.D., M.Bsc., FAAFP, at: Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, MD 20850, or by email to epc@ahrq.hhs.gov.

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# Quality Improvement Interventions To Address Health Disparities

Closing the Quality Gap: Revisiting the State of the Science

#### **Structured Abstract**

**Objective.** This review evaluates the effectiveness of quality improvement (QI) strategies in reducing disparities in health and health care.

**Data Sources.** We identified papers published in English between 1983 and 2011 from the MEDLINE<sup>®</sup> database, the Cumulative Index of Nursing and Allied Health Literature (CINAHL), Web of Science Social Science Index, and PsycINFO.

Review Methods. All abstracts and full-text articles were dually reviewed. Studies were eligible if they reported data on effectiveness of QI interventions on processes or health outcomes in the United States such that the impact on a health disparity could be measured. The review focused on the following clinical conditions: breast cancer, colorectal cancer, diabetes, heart failure, hypertension, coronary artery disease, asthma, major depressive disorder, cystic fibrosis, pneumonia, pregnancy, and end-stage renal disease. It assessed health disparities associated with race or ethnicity, socioeconomic status, insurance status, sexual orientation, health literacy/numeracy, and language barrier. We evaluated the risk of bias of individual studies and the overall strength of the body of evidence based on risk of bias, consistency, directness, and precision.

**Results.** Nineteen papers, representing 14 primary research studies, met criteria for inclusion. All but one of the studies incorporated multiple components into their QI approach. Patient education was part of most interventions (12 of 14), although the specific approach differed substantially across the studies. Ten of the studies incorporated self-management; this would include, for example, teaching individuals with diabetes to check their blood sugar regularly. Most (8 of 14) included some sort of provider education, which may have focused on the clinical issue or on raising awareness about disparities affecting the target population. Studies evaluated the effect of these strategies on disparities in the prevention or treatment of breast or colorectal cancer, cardiovascular disease, depression, or diabetes. Overall, QI interventions were not shown to reduce disparities. Most studies have focused on racial or ethnic disparities, with some targeted interventions demonstrating greater effect in racial minorities—specifically, supporting individuals in tracking their blood pressure at home to reduce blood pressure and collaborative care to improve depression care. In one study, the effect of a language-concordant breast cancer screening intervention was helpful in promoting mammography in Spanish-speaking women. For some depression care outcomes, the collaborative care model was more effective in lesseducated individuals than in those with more education and in women than in men.

**Conclusions.** The literature on QI interventions generally and their ability to improve health and health care is large. Whether those interventions are effective at reducing disparities remains unclear. This report should not be construed to assess the general effectiveness of QI in the health care setting; rather, QI has not been shown specifically to reduce known disparities in health care or health outcomes. In a few instances, some increased effect is seen in

disadvantaged populations; these studies should be replicated and the interventions studied further as having potential to address disparities.				

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# **Executive Summary**

# **Background**

Health care disparities are the differences or gaps in care experienced by one population compared with another. Disparities have been noted in health outcomes, including clinical outcomes such as mortality, process measures in the health care system, and disease prevalence. By definition, a disparity in health care quality or health outcomes is not due to differences in the health care needs or preferences of the patient but to other factors. Such differences in health outcomes and their determinants are associated with certain social conditions and demographic attributes. 3,4

Disparities that occur between identified populations are described by attributes such as race, ethnicity, language, sex, insurance status, socioeconomic status, and health literacy. These attributes and the disparities that may be associated with them are not mutually exclusive, and populations with disproportionately poor health outcomes often share multiple indicators of disparity. Despite what is known about disparities, it is not clear what strategies have the potential to improve the quality of care effectively and to reduce inequities for segments of the population.<sup>2</sup>

Quality improvement (QI) is a multidisciplinary, systems-focused, data-driven method of understanding and improving the efficiency, effectiveness, and reliability of health processes and outcomes of care. The QI process is designed to raise the standards of the delivery of preventive, diagnostic, therapeutic, and rehabilitative measures to maintain, restore, or improve the health outcomes of individuals and populations. Given the potential for QI strategies to improve the quality of care across the population, interest has developed in whether they might be used to reduce specific disparities, potentially by having an amplified effect among disadvantaged groups.<sup>5</sup>

For this report, we defined a QI intervention as a change process in health care systems, services, or suppliers for the purpose of increasing the likelihood of optimal clinical quality of care, measured by positive health outcomes for individuals and populations. An intervention could also be described as a strategy aimed at reducing the quality gap (the difference between health care processes or outcomes observed in practice and those potentially obtainable based on current evidence-based knowledge) for a group of patients representative of those encountered in routine practice.<sup>5</sup>

## **Objective**

This review evaluates the effectiveness of QI interventions in reducing disparities in health and health care.

# **Key Questions**

Key Question 1. What evidence is available about the effectiveness of quality improvement strategies to reduce differences in health outcomes associated with selected disparities in patients with key conditions?

Key Question 2. What evidence is available about the harms related to quality improvement strategies to reduce differences in health outcomes associated with selected disparities in patients with key conditions?

# **Analytic Framework**

We developed the analytic framework (shown in Figure 1 of the full report) based on clinical expertise and refined it with input from a Technical Expert Panel (TEP). The analytic framework outlines the review of the available evidence on the effectiveness of QI strategies in the reduction of disparities in health outcomes and other measures of health care delivery for selected conditions and groups.

We explicitly defined eligibility criteria using a PICOTS (population, intervention, comparator, outcome, timing, and setting) structure. Broadly, we sought studies that described a QI intervention and measured potential changes in the inequity of care between patient groups with prespecified clinical conditions.

To measure potential changes in disparity between patient groups, studies had to include a target and referent population (e.g., for income disparity studies, they should include data for low- and high-income groups). We included studies that reported outcomes in terms of health care processes, individual health outcomes, and/or adverse outcomes or harms resulting from a QI intervention.

#### **Methods**

## **Input From Stakeholders**

With input from our TEP, we drafted initial Key Questions (KQs), which were reviewed by the Agency for Healthcare Research and Quality. Our TEP also provided input during the project on issues such as setting, inclusion/exclusion criteria, and refining the analytic framework.

#### Literature Search

We searched the following databases: MEDLINE® (PubMed® interface), the Cumulative Index of Nursing and Allied Health Literature (CINAHL), Web of Science Social Science Index, and PsycINFO (CSA Illumina interface). The search strategies for each of these databases included terms related to QI, disparity, and prespecified clinical conditions. <sup>6,7</sup>

Each search strategy used a combination of subject headings (i.e., controlled vocabulary) and keywords. (See Appendix A of the full report.) We carried out hand searches of the reference lists of recent systematic reviews related to QI studies and the reference lists of included papers.

## **Paper Selection Process**

We included studies that captured health outcome measures and/or process measures to answer KQ 1. For KQ 2, we sought studies that reported harms (e.g., negative unintended consequences, misallocation of effort, decreased patient satisfaction) of the QI intervention to individual participants. Table A summarizes the inclusion/exclusion criteria.

Table A. Inclusion and exclusion criteria

Criteria		
Individuals receiving health care in the United States for a prespecified clinical condition:		
<ul> <li>Asthma</li> <li>Cancer: <ul> <li>Colorectal cancer (including screening)</li> <li>Breast cancer (including screening)</li> </ul> </li> <li>Cardiovascular disease <ul> <li>Congestive heart failure</li> <li>Coronary artery disease (including ischemic heart disease, myocardial infarction, and acute coronary syndrome)</li> <li>Hypertension</li> </ul> </li> <li>Cystic fibrosis</li> <li>Depression (major depressive disorder only)</li> <li>Diabetes</li> <li>End-stage renal disease</li> <li>Pneumonia (including pneumococcal vaccination)</li> <li>Pregnancy</li> </ul>		
Studies had to include data on characteristics known to be associated with health disparities: race or ethnicity, socioeconomic status, insurance status, sex, sexual orientation, health literacy/numeracy, and/or language barrier.		
QI strategy: (1) a formal broad organizational model or (2) a change process in health care systems, services, or suppliers for the purpose of increasing the likelihood of optimal clinical qualit of care.		
Usual care or use of an alternate strategy.		
Outcome measures of interest: health outcomes (e.g., morbidity and mortality, indirect health outcomes such as blood pressure and HbA1c); process measures (e.g., proportion of patients treated according to clinical guidelines); changes in disparity; and harms (i.e., any negative impact of the intervention on the individual patients or the health care system).		
1983-present		
Studies were based out of a hospital, provider office, and/or health care clinic.		
<ul> <li>Admissible designs: randomized controlled trials, including cluster randomized controlled trials; controlled trials, including quasi-randomized trials; controlled before-after studies; prospective and retrospective cohort studies; interrupted time series studies with comparison groups; and stepped-wedge design studies.</li> <li>Original research studies with sufficient detail to enable use and adjustment of the data and results.</li> <li>Inclusion of a target group and an internal or external referent group to measure changes in disparities.</li> <li>A minimum sample size of 50 individuals per study and intervention group or subgroup.</li> <li>Extractable data on relevant outcomes from text or tables.</li> <li>English-language publications only.</li> </ul>		

In the absence of published information (e.g., minimum effect size, standard error) to inform a power calculation, we derived the minimum sample size from expert opinion.

**Abbreviations:** HbA1c = hemoglobin A1c; QI = quality improvement.

As health care systems, disparities, and groups subject to disparities vary geographically, we limited eligible papers to studies of patients in the U.S. health care system. Consistent with this inclusion criterion, only papers published in English were included. Searches were limited to papers published in 1983 or later, as seminal work regarding QI strategies began to be published in the early 1980s.

All studies were required to include a comparison group that did not receive the QI intervention or that received a different intervention. In addition, they were required to provide data that could be used to measure a disparity before and after the intervention based on one of the population characteristics specified in the protocol (Table A). These data could have included reference to an external referent group, but if so, the data needed to have been collected within 4 years of the enrollment of the target group and be from a source that was at the State or local level. We included randomized controlled trials (RCTs), including cluster randomized controlled trials; controlled trials, including quasi-randomized studies; controlled before-after studies; prospective and retrospective cohort studies; interrupted time series with comparison groups; and stepped-wedge designs.

We considered both formal QI models and QI strategies for the review. We did not include papers describing topics or interventions covered by other reports in the Closing the Quality Gap series (e.g., studies that target public reporting, payment bundling, and medication adherence).

We conducted screening in two phases: abstract and full-text screening. Two reviewers independently reviewed each abstract. All papers with inclusion or exclusion conflicts at the abstract review level or lacking adequate information to make a determination were promoted to full-text review. Two reviewers independently reviewed the full text of papers included at the abstract phase. Disagreements between reviewers at the full-text screening level were resolved by a senior investigator.

#### **Data Extraction**

Two reviewers independently extracted relevant data (e.g., setting, condition, patient population, QI strategy, outcomes, and disparity) from all included papers using a predefined evidence table shell. A senior investigator reviewed the evidence tables for accuracy and completeness. The research team met regularly during the data extraction period and discussed global issues related to the process. The final evidence table is presented in Appendix I of the full report. When possible to identify, analyses resulting from the same study were grouped together.

# **Quality Assessment**

We assessed the quality of individual studies using specific tools for each type of study. For RCTs, we used the Cochrane Collaboration Risk of Bias tool, which evaluates domains that include sequence generation, allocation concealment, blinding, outcome data reporting, and reporting bias.

For observational studies, we used the Newcastle-Ottawa scale<sup>9</sup> to assess three broad perspectives: (1) the selection of the study groups, (2) the comparability of the groups, and (3) the ascertainment of either the exposure for case-control studies or the outcome of interest for cohort studies.

We rated individual studies as good, fair, or poor quality. Several of the included papers reported data from a post hoc or secondary analysis of a previously completed RCT. Because the balance between groups achieved by randomization does not reliably extend to subgroups, we modified the risk of bias/quality assessment on a case-by-case basis, considering the methods of the individual paper and parent study methods when appropriate.

## **Data Synthesis**

Meta-analysis was not appropriate in this review due to the heterogeneity of the studies in population, clinical condition, disparity target, and outcome; therefore, all analysis is narrative and based on the evidence and summary tables. Studies are summarized in categories of clinical conditions, and where possible, by type of outcome studied (e.g., clinical or process).

#### **Results**

## **Literature Search Yield**

Searches identified 4,278 titles and abstracts for screening. From this broad screening, 791 papers were identified as possibly related to our review and moved forward for full-text review. Nineteen papers met criteria; they represented 14 studies of cancer, cardiovascular disease, depression, and diabetes. All 14 studies included in the review addressed KQ 1, and none addressed KQ 2 (harms of interventions).

Of the 14 studies represented in the 19 included papers, 11 were RCTs, <sup>10-21</sup> including 2 cluster RCTs. <sup>10,11,22-25</sup> The remaining studies were cohort studies, including one prospective cohort study, <sup>26</sup> one retrospective cohort study, <sup>27</sup> and one cohort study with a historical control. <sup>28</sup>

Included papers targeted or described disparities associated with differences in race or ethnicity (n = 14),  $^{10-12,16,17,19-21,23,24,26-29}$  socioeconomic status (n = 3),  $^{13,19,27}$  insurance status (n = 2),  $^{14,21}$  language (n = 2),  $^{15,21}$  health literacy (n = 1),  $^{18}$  and sex (n = 1).  $^{25}$ 

Outcomes included health care processes and health outcomes. All but one of the studies incorporated multiple components into their QI approach. Patient education was a part of most interventions (12 of 14), although the specific approach differed substantially across the studies. Ten of the studies incorporated self-management—for example, teaching individuals with diabetes to check their blood sugar regularly. Most (8 of 14) included some sort of provider education, which may have focused on the clinical issue or on raising awareness about disparities affecting the target population. Nonetheless, given the degree to which the interventions all included multiple components that were implemented as a system, it is not possible to tease apart the effects or implications of individual aspects.

We organize the results in two ways. First, the results are summarized by effect on particular disparities, which is the primary focus of the review; second, descriptions of the studies are organized by clinical condition as a reference for end users interested in implementing QI approaches in individual clinics or clinical specialties.

# **Effects by Type of Disparity**

Eleven studies provided data on the effects of QI interventions on racial or ethnic disparities in health care (Table B). Among interventions to reduce racial or ethnic disparities, one disease management and patient education program<sup>27</sup> was associated with a reduction in disparity between Black and White patients in HbA1c (hemoglobin A1c) testing when it was targeted in a geographic area with very high rates of diabetes. This study reported significant improvement among Black participants compared with no improvement among White participants, thus narrowing the gap. Other interventions did not demonstrate a significant reduction in disparity but demonstrated an amplified effect in the nonwhite populations. They included an additional patient education program for reduction in blood pressures<sup>16</sup> and a complex collaborative care model aimed at providers of patients with depression. <sup>10,23,24,29</sup> In the latter study, the intervention

was more effective in the short term among minorities than among Whites, although the interaction was no longer significant after 1 year and the intervention was not effective overall at 5 and 9 years.

Table B. Summary of effects on disparities in health outcomes associated with race or ethnicity

Author, Year, Study Design, Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Arean et al., 2005 <sup>12</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>No disparity in depression severity existed by race or income at baseline.</li> <li>The intervention was effective in all racial subgroups, with no interaction by race and no amplified effect in any group.</li> <li>In subgroup analysis, the intervention was associated with greater use of psychotherapy but not pharmacotherapy within the Black population.</li> </ul>
Bao et al., 2011 <sup>19</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Provider reminder system</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>At baseline within the usual-care group, 22% of minorities had adequate antidepressant use, compared with 39% of Whites.</li> <li>The intervention had no effect on this disparity, and ethnic minorities did not receive greater benefit from intervention compared with Whites during any time period.</li> </ul>
Bosworth et al., 2011 <sup>16</sup> RCT Cardiovascular disease: hypertension	<ul> <li>Patient education</li> <li>Promotion of self-management</li> </ul>	<ul> <li>The race by time by treatment group effect model suggested differential intervention effects on BP over time for Whites vs. nonwhites for both SBP (p = 0.08) and DBP (p = 0.01).</li> <li>Compared with usual care, the combination of home BP monitoring and tailored behavioral intervention continued to be effective in nonwhite participants at 24 months (p = 0.04).</li> </ul>
Coberley et al., 2007 <sup>27</sup> Retrospective cohort Diabetes	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Organizational change (disease management)</li> </ul>	<ul> <li>Initial racial disparity in HbA1c testing between the diabetes HDZ group (higher than expected prevalence of diabetes) and non-HDZ group was 12%.</li> <li>Disparity was not significantly reduced after 12 months (p = 0.06).</li> <li>Within the HDZ zone, testing increased by 15% among Black participants but not among White participants, resulting in a reduction in disparity in this subgroup analysis.</li> </ul>

Table B. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Connett and Stamler, 1984 <sup>17</sup> RCT Cardiovascular disease: coronary artery disease and hypertension	<ul> <li>Patient education</li> <li>Promotion of self-management</li> </ul>	<ul> <li>At baseline, Black participants had higher rates of smoking than White participants (68.7% vs. 63%; p &lt; .001).</li> <li>Both racial groups experienced significant reductions in smoking, close to 50% in the intervention group and more than 35% in the usual-care group.</li> <li>The baseline disparity persisted in the intervention group but was apparently reduced in the usual-care group.</li> <li>A statistically significant but clinically insignificant disparity in DBP and SBP by race was present at baseline.</li> <li>Blood pressures were reduced in both the intervention and control groups, with greater change observed in the intervention group.</li> <li>The small disparity observed at baseline was further reduced at followup in the intervention group but not the control group.</li> </ul>
Lasser et al., 2011 <sup>21</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates existed at baseline by race or ethnicity.</li> <li>The intervention was more effective in White and Black individuals relative to those of other or unknown race.</li> </ul>
Mahotiere et al., 2006 <sup>26</sup> Prospective cohort Diabetes	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Other (community intervention)</li> </ul>	<ul> <li>The disparity in biennial lipid profile testing at baseline was 19%.</li> <li>The biennial lipid profile testing rate improved by 26.2% in African-American fee-for-service Medicare beneficiaries with diabetes in the intervention areas following implementation of the QI program.</li> <li>The disparity in performance of biennial lipid profile between African-American and White Medicare fee-for-service beneficiaries was reduced to 9.2% following implementation of the QI program.</li> <li>An analysis of the direct impact of the selected interventions on reducing the disparity in this uncontrolled database analysis was not feasible.</li> </ul>

Table B. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

(continued)		
Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Miranda et al., 2003; <sup>10</sup> Miranda et al., 2004; <sup>29</sup> Wells et al., 2007; <sup>23</sup> Wells et al., 2004 <sup>24</sup> Cluster RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>The intervention was associated with decreases in probable depressive disorder among minorities but not White patients at 12 months (Latino, p = 0.02; African-American, p = 0.01).</li> <li>At 12 months, among intervention recipients, the baseline disparity had increased from 6.7% to 7.7% between Latino and White patients and decreased from 9.2% to 6.7% between African-American and White patients.</li> <li>Although a statistically significant interaction was seen between intervention and ethnicity at 6 months when minorities were grouped and contrasted with White patients, no such interaction persisted at 12 months.</li> <li>The overall effect of the intervention on depression status was not significant at 5 and 9 years, but an interaction with race was seen in the overall model of effectiveness. The intervention was associated with improvements in the Mental Health Inventory among minorities (p = 0.008) but not among White patients (p = 0.59).</li> <li>In subanalysis at 5 years, QI-Therapy but not QI-Meds was effective within the minority population.</li> </ul>
Olomu et al., 2010 <sup>28</sup> Retrospective cohort (historic controls) Cardiovascular disease: coronary artery disease	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (guideline adherence)</li> </ul>	<ul> <li>The American College of Cardiology's Acute Myocardial Infarction Guidelines Applied in Practice strategy was associated with increased inpatient use of beta-blockers among nonwhite patients.</li> <li>Racial disparities in the use of cardiac catheterization and percutaneous coronary intervention appeared after implementation of the GAP QI strategy despite overall improvements in care.</li> <li>The admission tool and inpatient aspirin were more often used post-GAP vs. pre-GAP in both White and nonwhite patients.</li> </ul>
Sequist et al., 2010 <sup>11</sup> Cluster RCT Diabetes	<ul><li>Provider education</li><li>Audit and feedback</li></ul>	<ul> <li>Disparities between Black and White patients were present at baseline in HbA1c levels, BP control, and LDL level.</li> <li>The intervention showed no effect overall in either racial group.</li> <li>The intervention did not reduce the disparity.</li> </ul>
Siddiqui et al., 2011 <sup>20</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates existed at baseline by race or ethnicity.</li> <li>No statistically significant difference in screening rates existed between Whites and African-Americans in the control group.</li> <li>When intervention groups were combined, the screening rate was significantly higher in Whites than African-Americans.</li> </ul>

**Abbreviations:** BP = blood pressure; CRC = colorectal cancer; DBP = diastolic blood pressure; GAP = American College of Cardiology's Acute Myocardial Infarction Guidelines Applied in Practice; HbA1c = hemoglobin A1c; HDZ = health disparity zone; LDL = low-density lipoprotein; QI = quality improvement; RCT = randomized controlled trial; SBP = systolic blood pressure.

Two studies examined a difference in outcomes associated with insurance status (Table C). In both studies, the intervention was equally successful at increasing cancer screening in publicly and privately insured participants. In the first study, a patient reminder system for breast cancer screening improved mammography rates in all women. In the second study, language-concordant assistance by a patient navigator who promoted self-management strategies, patient education, and reminders were associated with significantly increased colorectal cancer screening among both privately insured and publicly insured participants compared with usual care but was more effective in the privately insured group.

Table C. Summary of effects on disparities in health outcomes associated with insurance status

Author, Year, Study Design, Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Barr et al., 2001 <sup>14</sup> RCT Cancer: breast cancer screening	Patient reminder system	<ul> <li>No disparity in mammography screening rates was observed at baseline.</li> <li>The intervention was successful in both groups.</li> <li>Reminder interventions improved the likelihood of screening mammography in both commercially insured women (p = 0.001) and women covered by Medicare (p = 0.01), with no difference in improvement between groups.</li> </ul>
Lasser et al., 2011 <sup>21</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates was measured at baseline by race or ethnicity.</li> <li>The intervention increased screening rates in both the private and public insurance groups compared with individuals in the usual-care group.</li> <li>The intervention was associated with a better screening rate for the privately insured group than the publicly insured group.</li> </ul>

**Abbreviations:** CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial.

Two studies examined the effects of QI strategies on disparities associated with language (Table D). Both of them studied language concordance, in which strategies are provided in the native or preferred language of the participant (e.g., in Spanish for native Spanish speakers). One study examined the degree to which a language-concordant patient education strategy was associated with increased cancer screening (breast and colorectal) among English- and Spanish-speaking patients. For breast cancer screening, Spanish speakers were more likely to be up to date at baseline than English speakers (odds ratio [OR], 1.46; 95% confidence interval [CI]: 1.16 to 1.84). The intervention was associated with increased rates of screening overall, with subgroup analysis indicating a greater effect in the Spanish-speaking group (OR, 1.85; 95% CI: 1.38 to 2.47) than the English-speaking group (OR, 1.18; 95% CI: 0.82 to 1.71). However, the overall multivariate analysis failed to confirm these results, and providing the intervention in Spanish to Spanish speakers did not make it any more effective in this group. For colorectal screening, there was no difference in up-to-date status at baseline, the intervention was again effective overall, and there was no language-by-intervention effect.

A second study included language-concordant assistance by a patient navigator promoting self-management strategies, and providing patient education and reminders to facilitate adherence to colorectal cancer screening for individuals speaking English as their primary language and individuals speaking a language other than English. The patient navigator intervention was associated with increased colorectal cancer screening among individuals whose

primary language was not English (28.9 percent vs. 18.9 percent; p = 0.04) but not among patients whose primary language was English (26.8 percent vs. 21.4 percent; p = 0.35). These studies combined may suggest that targeted language-concordant interventions could warrant further examination, with results suggesting a significantly different effect for non-English speakers and English speakers in one study, and a clinically but not statistically different effect in the other.

Table D. Summary of effects on disparities in health outcomes associated with language barrier

Author, Year, Study Design, Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Beach et al., 2007 <sup>15</sup> RCT Cancer: CRC and breast cancer screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>At baseline, Spanish speakers were more likely to be up to date on breast cancer screening.</li> <li>The intervention was effective at increasing rates of breast cancer screening overall, with greater effect among Spanish speakers.</li> <li>The difference between observed effects for breast cancer screening in the two language groups was not significant.</li> <li>No disparity in CRC screening rate was observed at baseline.</li> <li>The intervention was associated with increases in CRC screening in both groups, with neither group having a greater effect of the intervention.</li> <li>Although there was no evidence that the intervention might reduce known disparities, the intervention was effective at increasing CRC screening for both groups.</li> </ul>
Lasser et al., 2011 <sup>21</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC was measured at baseline.</li> <li>English-speaking participants had a similar incidence of CRC screening during 1 year of followup in the intervention group as compared with usual care.</li> <li>Intervention was particularly beneficial for non-English-language participants.</li> </ul>

**Abbreviations:** CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial.

In one study focused on improving provider-patient communication in Department of Veterans Affairs clinics, colorectal cancer screening increased among individuals with limited health literacy (55.7 percent vs. 30 percent) but not among individuals with adequate health literacy (39 percent vs. 36 percent) in the 20-percent subsample that underwent literacy assessment (Table E). The intervention itself included a workshop and feedback sessions for providers and educational materials for patients that included a video.

Table E. Summary of effects on disparities in health outcomes associated with health literacy

Author, Year, Study Design, Clinical Condition  QI Interv	Ettect on Health Disparity
Ferreira et al., 2005 <sup>18</sup> RCT  Cancer: CRC screening  • Patient et en provider et al., 2005 • Audit and	ducation more likely to be screened for CRC when treated at

Abbreviations: CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial; VA = Veterans Affairs.

In the two studies that assessed differences in effect by socioeconomic status, no effect was seen by income, but individuals with less education experienced greater benefits of collaborative care for depression than did those with higher education (Table F).

Table F. Summary of effects on disparities in health outcomes associated with socioeconomic status

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Author, Year, Study Design, Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Arean et al., 2007 <sup>13</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	Both low-income populations and those with high/middle income experienced a very small benefit from the collaborative care intervention: fewer depression symptoms (adjusted OR, -0.41; 95% CI: -0.49 to -0.33 for high/middle income; adjusted OR, -0.39; 95% CI: -0.5 to -0.27 for low income; comparator: usual care). However, no disparities in depressive symptoms had existed at baseline.
Bao et al., 2011 <sup>19</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Provider reminder system</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>No disparity in depressive symptoms was present at baseline.</li> <li>At 24 months, participants with no college education had a greater reduction in depression than participants with college education.</li> </ul>

**Abbreviations:** CI = confidence interval; OR = odds ratio; QI = quality improvement; RCT = randomized controlled trial.

Finally, one analysis examined the degree to which a collaborative care model for depression could reduce known disparities by sex in accessing care and in outcomes (Table G). At baseline, women were more likely to have current single or double depression (62 percent) than men (53 percent) and had more symptoms of depression and lower mental health–related quality of life. Women had higher rates of appropriate depression care compared with men at 2 years (p = 0.0001). A medication-focused intervention and a therapy-focused intervention decreased a

disparity gap between men and women in probable unmet need from 10 percent to 1 percent (QI–Meds) and 3 percent (QI–Therapy) at 24 months.

Table G. Summary of effects on disparities in health outcomes associated with sex

Author, Year, Study Design, Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Sherbourne et al., 2004 <sup>25</sup> Cluster RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>At baseline, women were more likely to have depression (62%) than men (53%) and had more depression symptoms and lower mental health–related quality of life.</li> <li>Women had higher rates of appropriate depression care compared with men at 2 years (p = 0.0001).</li> <li>QI–Meds and QI–Therapy decreased "probable unmet need" disparity gap between men and women from 10% to 1% (QI–Meds) and 3% (QI–Therapy) at 24 months.</li> </ul>

**Abbreviations:** QI = quality improvement; RCT = randomized controlled trial.

# **Studies by Clinical Condition**

#### Cancer

Five RCTs with subgroup analyses explored the effects of various QI strategies on health care disparities in cancer screening, including one examining breast cancer screening, <sup>14</sup> three assessing screening for colorectal cancer, <sup>18,20,21</sup> and one assessing both breast and colorectal cancer screening. <sup>15</sup> Disparities that served as the focus of these analyses included race or ethnicity, <sup>20,21</sup> insurance status, <sup>14,21</sup> health literacy, <sup>18</sup> and language. <sup>15,21</sup>

The QI strategies included provision of mail or telephone reminders to patients, <sup>14</sup> education and feedback for clinicians, <sup>18</sup> and language-concordant telephone support calls from prevention care managers to patients. <sup>15</sup> All five RCTs took place in the United States, with study settings including a large academic medical center, <sup>20</sup> a primary care research network, <sup>21</sup> 1 large group-model health maintenance organization (HMO), <sup>14</sup> 2 Department of Veterans Affairs (VA) clinics, <sup>18</sup> and 11 community health centers. <sup>15</sup>

All studies employed an internal usual-care comparison group. Compared with usual care, a language-concordant intervention was more effective in increasing breast cancer screening among Spanish-speaking women than English-speaking women, but the observed difference between the two groups (English and Spanish speaking) was not significant. The language-concordant intervention did not have a similar effect on colorectal cancer screening. Compared with usual care, a strategy targeting health literacy facilitated colorectal cancer screening among those with limited health literacy more effectively than among those with high health literacy. A reminder intervention for breast cancer screening had no differential effect on mammography disparities by insurance status.

#### **Cardiovascular Disease**

One post hoc analysis of an RCT<sup>17</sup> and one retrospective cohort study<sup>28</sup> explored the effects of various QI strategies on racial health care disparities in coronary artery disease (CAD). The RCT addressed reduction of CAD risk factors,<sup>17</sup> while the retrospective cohort examined management of acute myocardial infarction (AMI).<sup>28</sup> QI strategies included patient education

and facilitation of self-management,<sup>17</sup> and a multifactorial provider- and systems-focused strategy.<sup>28</sup> Both studies were collaborations of academic and community health centers.<sup>17,28</sup> The studies each employed an internal usual-care comparison group.

One study of cardiovascular risk factor modification showed no meaningful reduction in health disparities seen in smoking rates, although both Black and White participants had substantially lower rates of smoking after intervention.<sup>17</sup> In the other study, intervention in AMI treatment reduced disparities in one aspect of treatment, which exacerbated disparities in other areas, including use of the discharge tool and cardiac catheterization rates.<sup>28</sup> The strength of evidence was insufficient.

Two post hoc analyses of RCTs explored the effects of various QI strategies on racial health care disparities in hypertension. The RCTs addressed management of hypertension and reduction of CAD risk factors, including hypertension. Ull strategies were patient education and facilitation of self-management. The studies took place in university clinics and multicenter collaborations of academic and community health centers. The studies each employed an internal usual-care comparison group.

One study had no significant intervention effect on a clinically insignificant disparity in blood pressure measures present at baseline after patient education and promotion of self-management.<sup>17</sup> In the second study, a home-based self-management strategy, including home blood pressure monitoring and tailored self-management strategies, was more effective in the Black population than in the White population, although the study design precludes determination of a clear causal effect from the intervention.<sup>16</sup>

## **Depression**

Three studies evaluated the effect of QI interventions on disparities in depression outcomes. Racial disparities were of interest in all three, but interim analyses were also performed based on sex, 25 income, 13 and educational status. 19 All three studies used a collaborative care model, which involved collaboration among multiple clinical providers to provide a coordinated set of interventions. The model in all three studies generally included a dedicated mental health coordinator (nurse or case manager); creation of mental health teams (composed of primary provider, facility nurses, and psychiatrists); evidence-based pharmacotherapy and psychotherapy; extensive provider education; and longitudinal patient followup to evaluate clinical status and adherence. Each intervention was designed to address known barriers to the receipt of quality mental health care. All three studies were prospective RCTs, with randomization occurring at the practice level and referring to training provided to the providers. However, individual providers and patients retained the ability to select the treatment provided to the individual patient. All three trials took place in the United States.

The collaborative care models described in this report were all associated with improvements in mental health outcomes, including depression scores, severity, and functioning, but none specifically demonstrated a reduction in disparity caused by the intervention. In part, this was because few disparities were measurable at baseline. The studies showed that there was no significant difference in the effect in groups defined by income, race, or education. Nonetheless, there were some notable differences in effectiveness that might inform future research. For example, one study demonstrated a greater effect on clinical outcomes in the less educated group, <sup>19</sup> and the effect of a second intervention was amplified in minorities on some measures. <sup>23</sup> Although no change in disparity was associated with the interventions, improvements occurred across the board, and no harms were reported in any of the studies.

#### **Diabetes**

Three good-quality studies assessed the effect of QI interventions on disparities in diabetes outcomes. One was an RCT,<sup>11</sup> one was a prospective cohort study,<sup>26</sup> and one was a retrospective cohort study.<sup>27</sup> All of these studies reported on surrogate clinical outcomes, clinical risk factors for diabetes comorbidities, and process measures. In two of three studies, disparities were reduced in one or more outcomes for at least one subgroup, but the study designs were such that the reduction could not be shown to be caused by the intervention.<sup>26,27</sup> In one study of a patient reminder system, racial disparities were reduced when HbA1c testing increased substantially among Black participants relative to no change among White participants. In a broad systems-level program in New York State, a disparity of 19 percent in biennial lipid testing between Black and White Medicare recipients was reduced to 9.2 percent after intervention of a QI program.

#### **Discussion**

We identified individual studies that suggest benefits in particular subgroups known to suffer from disparities in health and health care, but evidence is unavailable to guide QI efforts specifically to reduce disparities. Although there is limited evidence available, several strategies are worthy of future study and possibly wider implementation. These strategies include the collaborative care model and targeted patient education, including language and literacy concordance. Data are insufficient to support universal implementation of these strategies, but the strategies may be suitable for implementation if an appropriate plan is in place to monitor their effectiveness and potential adverse effects.

Most studies have focused on racial or ethnic disparities. Some targeted interventions have demonstrated greater effect in racial minorities: specifically, supporting individuals in tracking their blood pressure at home to reduce blood pressure and collaborative care to improve depression care. Language concordance was evaluated in only one study, but a language-concordant breast cancer screening intervention was helpful in promoting mammography in Spanish-speaking women. The collaborative care model in depression was more effective in less educated individuals than in those with more education, and was more effective in women than in men for some depression care outcomes. None of the evidence is adequate to be confirmatory, but these studies suggest areas for future evaluation and targeted approaches.

Despite positive results seen in specific studies on specific clinical outcomes in some or all study populations, the strength of the evidence for QI interventions reviewed in this report *to affect disparities* is insufficient. Although adequate evidence exists from other sources to suggest the benefit of QI interventions in improving outcomes for a clinical population, the degree to which these interventions might be used to close an existing disparity gap has not been clearly demonstrated.

Our assessment is consistent with at least one prior review (from 2006),<sup>30</sup> and despite a larger body of literature on QI today and the presence of research demonstrating the effectiveness of QI interventions across populations, evidence for the effects of QI interventions on gaps in care related to disparities remains limited. Few studies focus specifically on reducing gaps in the availability, accessibility, and quality of health care between any two populations. Authors of studies in this review have attempted to address the question by conducting post hoc analyses of RCTs intended to study the effectiveness of QI interventions; however, in doing so, they have broken what randomization existed and have been unable to make the comparison necessary to tie observed improvements to the QI intervention conclusively.

One of the challenges in conducting a systematic review of the degree to which QI interventions can address disparities is the substantial breadth and heterogeneity of clinical conditions of interest, populations with the clinical conditions, QI intervention strategies, comparators, important clinical outcomes, surrogate outcomes, and disparities of interest. Compounding this heterogeneity are challenges to indexing QI strategies in the medical literature databases. For example, the subject term "Quality Improvement" was added to the National Library of Medicine's Medical Subject Heading Database (MeSH) only in 2011; before this time, myriad subject terms were used to index the various strategies described by authors of the QI literature, understandably leading to tremendous variability in how similar studies are categorized in the database. This partially reflects a lack of consistency about what constitutes a QI intervention; information on QI interventions available in the literature is often not clearly identified as such, and interventions may be multifaceted and thus difficult to evaluate or compare with other interventions. Many studies identified in the literature as including QI interventions also include non-QI interventions, such as broader public health initiatives; thus, the potential impact of the QI intervention may be masked or difficult to isolate.

Further challenges to studying changes in disparities are the poor documentation of disparities and the fact that many individuals experience multiple and overlapping disparities. Many of the studies we found that might have been able to empirically assess a disparity change were unable to demonstrate any existing disparity at baseline. Future studies will require much broader populations that include enough individuals from diverse backgrounds to capture and assess disparities over time empirically.

# **Applicability**

Although we reviewed fairly large studies conducted in diverse areas of the United States, all of the studies had substantial gaps in applicability to one or more populations of patients likely to present with the condition under study. Therefore, health systems or clinicians wishing to replicate any of these interventions should carefully assess whether the interventions apply or must be modified to suit their particular patient population, clinical setting, and available resources.

The overall insufficient strength of evidence suggests that decisions about whether to replicate interventions in this study and under what circumstances they should be replicated must be made without confidence in the degree to which disparities might be narrowed. By far the largest proportion of the literature focused on the ability of QI interventions to reduce racial disparities, with some suggestions that targeted programs could have some greater effects among racial minorities in both diabetes<sup>27</sup> and hypertension. <sup>16</sup> Far less information is available about QI interventions targeting other disparities, and the degree to which available evidence is applicable to other clinical conditions, other disparities, and other interventions is an area of potentially rich research. Health systems and individuals wishing to apply QI strategies are likely to be concerned about their applicability within clinical conditions, given the structure of the health system. Therefore, we summarize applicability by clinical focus below.

#### Cancer

Studies included patients cared for at community clinics in New York City, men treated at two VA clinics in Chicago, and women enrolled in a large group-model HMO in the northeastern United States. These settings were appropriate for cancer screening interventions, as the bulk of

cancer screening recommendations focus on the clinic setting. However, it is uncertain how well the results of these studies can be generalized to other populations or settings.

The tested interventions varied substantially, ranging from patient reminders to provider education with audit and feedback. These interventions could be replicated, although they generally required significant organizational resources to develop and implement and may not be feasible in other settings. Barriers to care may also differ in other settings, and the interventions likely would need to be adapted to the needs of the target population. In each study, usual care served as the comparator, and this too may differ in other practice settings. Thus, the marginal benefit of each intervention likely would be different in different settings.

Study outcomes consisted only of short-term process measures (i.e., receipt of cancer screening during followup). No long-term outcomes or clinical outcomes, such as diagnosis of malignancies, were reported. Thus the long-term clinical impact of such interventions is unclear.

#### **Cardiovascular Disease**

Studies of CAD risk factor control included men with CAD risk factors at clinical centers in 18 U.S. cities and patients with hypertension cared for at two university-affiliated clinics in North Carolina. A study involving AMI treatment included patients hospitalized at academic and community hospitals in Michigan. The study involving men only has limited applicability to women, as patterns of CAD risk factors differ by sex. Moreover, its enrollment occurred between 1973 and 1975, limiting applicability to present-day practice. Of the other two studies, one's results are applicable to patients in academic primary care practices, and the other's results are applicable to academic or community hospitals.

The interventions for CAD risk factor control included intensive patient education and self-management, along with medication titration in one study. The intervention for AMI treatment involved provider education, practice feedback, and implementation of a toolkit. These all required significant institutional resources, and the CAD risk factor interventions in particular may not be feasible in routine clinical practice. The AMI treatment initiative, although requiring institutional commitment, has already been disseminated extensively around the United States as a professional society initiative (American College of Cardiology Guidelines Applied in Practice); thus, its replication is confirmed to be feasible. In each of these studies, usual care served as the comparator. As this varies across practice settings, the effect of the interventions may differ in other environments.

For studies of cardiovascular risk factor control, outcomes consisted of intermediate clinical variables (hypertension, cholesterol, smoking, weight). Outcome assessment in the AMI treatment study was extensive but focused on measures of process and proximal utilization (e.g., prescription of evidence-based medications, use of cardiac catheterization).

## **Depression**

Two of the three studies focused on elderly patients in primary care. One included a range of ages in adulthood. All included both men and women and were racially diverse. Nonetheless, these patient groups may represent a small proportion of the individuals who struggle with depression because of the limited range of health care settings represented in these studies. It is unclear whether the observed results apply to patient populations who receive their primary and mental health care outside of a managed care system or to individuals who do not receive regular medical care. Additionally, given the settings in which the studies took place, they also may not apply to vulnerable populations receiving care through public health systems.

The interventions were all intensive in terms of demand on resources and required strong communication between care providers. In one study, enrolled practices committed to an intervention cost-sharing arrangement, with the understanding that the long-term implementation would fall on the organization of practice itself. The degree to which this is likely to be feasible is unclear.

All of the studies compared the intervention with usual care, although usual care was not ever completely described and therefore would be expected to vary.

Generally speaking, outcome measures were appropriate and reflected those that would and could be used in practice. They included changes in depressive symptoms, incidence of probably depressive disorder, mental health–related quality of life, functional impairment, and receipt of appropriate depression care.

All of the studies were conducted in primary care practices associated with larger health care organizations. It is unclear whether results would apply to other settings, including individual practices without the resources of a larger organization or assisted living facilities (pertinent because of the focus on the elderly population).

#### **Diabetes**

Studies included people cared for by primary care clinicians in ambulatory health centers in eastern Massachusetts, diabetes disease management program members living in socioeconomically disparate areas throughout the United States, and Medicare patients in New York State. The results may or may not be applicable to other populations in other regions.

Interventions evaluated included cultural competency training for clinicians and race-stratified performance reports with recommendations for Black patients with diabetes, patient telephone reminders in health disparity zones (defined as areas with diabetes prevalence above the national average for minorities), and Medicare New York State Quality Improvement Organization (IPRO) multifaceted provider and community interventions. The interventions may not be available in other regions and settings, since they required significant programmatic and implementation resources. The usual-care comparators described in these studies may not be applicable to other settings and regions.

Studies reported surrogate clinical outcomes (i.e., HbA1c control), clinical risk factors for diabetes comorbidities (i.e., blood pressure and lipid control), and process measures (i.e., HbA1c and low-density lipoprotein measurements). Duration of studies was generally 1 year. No studies reported any critically important clinical outcomes of diabetes, such as death or microvascular and/or macrovascular complications. Results from surrogate outcomes may not apply to important long-term clinical outcomes in people with diabetes.

Studies were conducted in ambulatory health centers in eastern Massachusetts, in diabetes disease management programs across the United States, and in New York State. As much diabetes care is delivered in primary care ambulatory settings, the evidence would be applicable. However, specialty clinic settings were not reported and the evidence may not apply to them.

#### **Conclusions**

The literature on QI interventions generally and their ability to improve health and health care is large. Whether those interventions are effective at reducing disparities remains unclear. This report should not be construed to assess the general effectiveness of QI in the health care setting; rather, QI has not been shown specifically to reduce known disparities in health care or health outcomes. In a few instances, some increased effect is seen in disadvantaged populations;

these studies should be replicated and the interventions studied further as having potential to address disparities.

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## Introduction

## **Background**

Health care disparities are the differences or gaps in care experienced by one population compared with another. Disparities have been noted in health outcomes including clinical outcomes and mortality, process measures in the health care system, and disease prevalence. By definition, a disparity in health care quality or health outcomes is not due to differences in health care needs or preferences of the patient, but to other factors. <sup>2</sup>

Disparities occur between identified populations, described by attributes such as race, ethnicity, language, sex, insurance status, socioeconomic status, and health literacy.<sup>2</sup> These attributes and the disparities that may be associated with them are not mutually exclusive, and populations with disproportionately poor health outcomes often share multiple indicators of disparity.

Examples of health disparities include substantially higher rates of death due to coronary heart disease among Black men and women than their White counterparts. Rates of preventable hospitalizations are inversely related to income. Hypertension is substantially more prevalent among Black than other populations, and tobacco use is higher among minorities, particularly Native Americans.<sup>3</sup>

Nonetheless, despite well documented disparities in health and health care in the United States, little is known about what interventions might serve to reduce differences. Given the potential for quality improvement (QI) efforts in the health care setting to improve outcomes in general, considering whether these interventions could be fruitful in reducing disparities as well is a logical next step.

# **Quality Improvement**

QI is a multidisciplinary, systems-focused, data-driven method of understanding and improving the efficiency, effectiveness, and reliability of health processes and outcomes of health care. The QI process is designed to raise the standards of the delivery of preventive, diagnostic, therapeutic, and rehabilitative measures to maintain, restore or improve health outcomes of individuals and populations. The ongoing process of QI requires all four of the following elements: performance goals, performance measures, QI practices, and feedback and reporting.<sup>4</sup>

QI describes a wide range of initiatives aimed at improving quality in health care organizations and includes programmed approaches that build on models and tools first used in industry as many as 60 years ago. One of the earliest approaches was Plan, Do, Study, Act (PDSA), and other examples include Total Quality Management (TQM), Continuous Quality Improvement (CQI), Business Process Reengineering (BPR), rapid cycle change, lean thinking, and Six Sigma. Another model is Focus, Analyze, Develop, Execute/Evaluate (FADE).

For this report, we defined a QI intervention as a change process in health care systems, services, or suppliers for the purpose of increasing the likelihood of optimal clinical quality of care, measured by positive health outcomes for individuals and populations. An intervention could also be described as a strategy aimed at reducing the quality gap (the difference between health care processes or outcomes observed in practice and those potentially obtainable based on current evidence-based knowledge) for a group of patients representative of those encountered in

routine practice.<sup>6</sup> As a starting point, we borrowed the taxonomy of QI strategies described in the antecedent Closing the Quality Gap series.<sup>6</sup> Examples of QI strategies that we expected to find in the literature are found in Table 1.

Table 1. Quality improvement strategies examples

QI Strategy	Examples	
Patient education	<ul> <li>Classes</li> <li>Parent and family education</li> <li>Patient pamphlets</li> <li>Intensive education strategies promoting self-management of chronic conditions</li> </ul>	
Provider education	<ul> <li>Workshops and conferences</li> <li>Educational outreach visits (e.g., academic detailing)</li> <li>Distribution of educational materials</li> </ul>	
Promotion of self- management	Materials and devices to promote self-management	
Audit and feedback	<ul> <li>Feedback of performance to individual providers</li> <li>Quality indicators and reports</li> <li>National/state quality report cards</li> <li>Publicly released performance data</li> <li>Benchmarking – provision of outcomes data from top performers for comparison with provider's own data</li> </ul>	
Facilitated relay of clinical data to providers	<ul> <li>Transmission of clinical data from outpatient specialty clinic to primary care provider by means other than medical record, (e.g., phone call or fax)</li> </ul>	
Patient reminder systems	Postcards or calls to patients	
Provider reminder systems	<ul> <li>Reminders in charts for providers</li> <li>Computer-based reminders for providers</li> <li>Computer-based decision support</li> </ul>	
Organizational change	<ul> <li>Case Management, Disease Management</li> <li>Total Quality Management, Cycles of Quality Improvement</li> <li>Multidisciplinary teams</li> <li>Change from paper to computer-based records</li> <li>Increased staffing</li> <li>Skill mix changes</li> </ul>	
Other	<ul> <li>Guideline adherence</li> <li>Care manager</li> <li>Collaborative care model</li> </ul>	

# **Disparity**

A health disparity is the difference between health care processes or outcomes observed in practice for a specific population, compared with another population. By definition, a disparity in health care quality or health outcomes is not due to differences in health care needs or preferences of the patient, but to other factors.2 Differences in health outcomes and their determinants between segments of the population are associated with certain social conditions, and demographic attributes.3,7

Disparities may be associated with attributes such as socioeconomic status (low income), minority group (race, ethnicity, culture, language), sex (women), age (children and elderly), access (insurance status), geography (inner-city or rural), education (health literacy), or disability. Numerous environmental or contextual differences, (e.g., resource availability,

transportation, air pollution) may also contribute to inequitable health care quality. As noted in the Centers for Disease Control and Prevention Health Disparities and Inequalities Report,3 the terms health disparity, health inequality, and health inequity are sometimes used interchangeably and each may provide an important indicator of community health.

Consistent with extensive research and findings in previous reports, the 2010 National Healthcare Disparity Report (NHDR) found that disparities related to race, ethnicity, and socioeconomic status are pervasive in the American health care system.1 Within the scope of health care delivery, these disparities may be due to differences in access to care, provider biases, poor provider-patient communication, poor health literacy, or other factors. Three key themes emerged from the report:

- 1. Disparities are common and uninsurance is an important contributor.
- 2. Many disparities are not decreasing over time.
- 3. Some disparities merit particular attention, especially disparities related to care for cancer, heart failure, and pneumonia.

The report also included a charge to examine disparities in "priority populations," which are groups with unique health care needs or issues that require special attention. The identification of priority populations has remained consistent across Institute of Medicine and Agency for Healthcare Research and Quality (AHRQ) and the Evidence-based Practice Center (EPC) Program documents from 2003 through 2010.

## **Approaches To Reducing Disparities**

Despite what is known about disparities, it is not clear what strategies have the potential to effectively improve the quality of care and to reduce inequities for segments of the population. QI interventions have been successful at improving health outcomes generally and in a number of settings; it is possible that they could be adapted or targeted to narrow a health or health care gap. If they were implemented in either a targeted way (i.e., in locations with especially high disparities) or broadly, they could potentially affect disparities at the population level.

A prior systematic review and analysis conducted by Beach and colleagues in 2006 summarized controlled studies of interventions targeted at health care providers to improve health care quality or reduce disparities in care for racial or ethnic minorities. Twenty-seven studies met criteria for review. Almost all (n = 26) took place in the primary care setting, and most (n = 19) focused on improving provision of preventive services.<sup>8</sup>

Nonetheless, the report concluded that there was little evidence to promote the use of QI interventions specifically to reduce disparities. The particular difficulty in assessing the impact of any health care intervention on disparities is that research must show effectiveness across multiple planes. Evidence of the effectiveness of the intervention needs to be demonstrated using a non-intervention comparison group, and at the same time, a disparity in outcome must be narrowed in the intervention group, but not in the comparison group. Thus, for a QI intervention to be effective for reduction of disparity both intervention effectiveness and disparity reduction effectiveness must be demonstrated; the intervention would be *more* effective for disadvantaged groups or individuals than for advantaged groups. A judgment of effectiveness in reducing disparities is therefore not possible when the intervention is targeted only at disadvantaged individuals or groups. While extraordinarily important, research that demonstrates a change in both effectiveness outcomes and disparity between at least two groups is thus complicated and rare.

# **Scope and Key Questions**

## **Scope of This Report**

The objective of this review was to assess the effectiveness of QI interventions in reducing disparities in health and health care.

## **Key Questions**

Key Question 1 (KQ 1). What evidence is available about the effectiveness of quality improvement strategies to reduce differences in health outcomes associated with selected disparities in patients with key conditions?

Key Question 2 (KQ 2). What evidence is available about the harms related to quality improvement strategies to reduce differences in health outcomes associated with selected disparities in patients with key conditions?

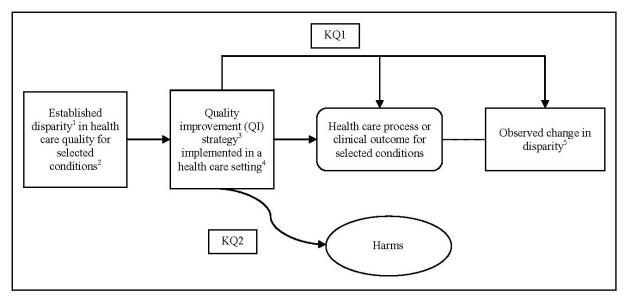
## **Analytic Framework**

We developed the analytic framework (Figure 1) based on clinical expertise and refined it with input from our TEP members. The analytic framework outlines the review of the available evidence on the effectiveness of QI strategies in the reduction of disparities in health outcomes and other measures of health care delivery for selected conditions and groups.

We explicitly defined eligibility criteria using a PICOTS (population, intervention, comparator, outcome, timing, and setting) structure (Table 2). Broadly, we sought studies that described a QI intervention and measured potential changes in the inequity of care between patient groups with pre-specified clinical conditions.

To measure potential changes in disparity between patient groups, studies had to include a target and referent population (e.g., for income disparity studies, they should include data for low and high income groups). We included studies that reported outcomes in terms of health care processes, individual health outcomes, and/or adverse outcomes or harms resulting from a QI intervention.

Figure 1. Analytic framework



**Abbreviations:** KQ = Key Question, QI = quality improvement.

<sup>1</sup>Disparities include: race or ethnicity, socioeconomic status, insurance status, sex, sexual orientation, health literacy/numeracy, and language barrier. <sup>2</sup>Selected conditions include: asthma, cardiovascular disease (including congestive heart failure, coronary heart disease, and hypertension); cancer (specifically colorectal cancer and breast cancer); cystic fibrosis; depression; diabetes; end stage renal disease; pneumonia (including pneumococcal vaccination); and pregnancy. <sup>3</sup>Taxonomy of quality improvement strategies: patient education; provider education; promotion of self-management; audit and feedback; facilitated relay of clinical data to providers; organizational change; patient reminder systems; and provider reminder systems. <sup>4</sup>Settings include those in which QI interventions were tested: hospitals, provider offices, and/or health care clinics. <sup>5</sup>Inclusion of a target and referent group is required to demonstrate a disparity.

## **Uses of This Report**

This evidence report addresses the KQs using the methods described to conduct a systematic review of published literature. We anticipate that the report will primarily be of value to researchers interested in studying disparities and to funders developing RFAs for this type of research. It may also be of value to policymakers and health systems leaders as they consider the potential impact of QI interventions focused on addressing disparities in health care.

## **Methods**

# **Topic Refinement**

Topics for the Closing the Quality Gap series were solicited from the portfolio leads at the Agency for Healthcare Research and Quality (AHRQ). The nominations included a brief background and context; the importance and/or rationale for the topic; the focus or population of interest; relevant outcomes; and references to recent or ongoing work.

Among the topics that were nominated, the following considerations were made in selection for inclusion in the series: the ability to focus and clarify the topic area appropriately; relevance to quality improvement (QI) and a systems approach; applicability to the Evidence-based Practice Center (EPC) program/amenable to systematic review; the potential for duplication and/or overlap with other known or ongoing work; relevance and potential impact in improving care; and fit of the topics as a whole in reflecting AHRQ portfolios.

Following assignment of the topic to the EPC, we identified technical experts on the topics of QI and disparities. The Technical Expert Panel (TEP) members contributed to AHRQ's broader goals of (1) creating and maintaining science partnerships as well as public-private partnerships and (2) meeting the needs of an array of potential customers and users of its products.

# **Literature Search Strategy**

#### **Databases**

We searched the following databases: MEDLINE (PubMed interface), the Cumulative Index of Nursing and Allied Health Literature (CINAHL), Web of Science Social Science Index, and PsycINFO (CSA Illumina interface). The search strategies for each of these databases included terms related to QI, disparity, and pre-specified clinical conditions. <sup>10,11</sup>

Each search strategy used a combination of subject headings (i.e., controlled vocabulary) and keywords (see Appendix A). We carried out hand searches of the reference lists of recent systematic reviews related to QI studies and the reference lists of included papers.

# **Study Selection**

# **Study Populations**

Eligible studies included individuals receiving health care in the U.S. Studies had to include data on one or more of the following characteristics known to be associated with health disparities: race or ethnicity, socioeconomic status, insurance status, sex, sexual orientation, health literacy/numeracy, and language barrier.

# **Sample Size**

As an inclusion criterion, we set the study sample size at a minimum of 50 participants each for the comparison and intervention groups, including subgroups that were stratified for analysis of intervention effect on disparity. The typical study designs used for QI studies have an inherent risk of bias. The maximum improvement possible in a QI trial is 100 percent; if the baseline performance is high, then the margin for improvement is so small that a high number of

participants would be required to rule out an observed effect due to chance. Because the conditions and outcomes were very diverse, it was not possible to forecast a single prevalence for the comparison group, nor a single minimum relative effect.

Setting a minimum sample size based on quantitative power calculations would have required studies in excess of 100 individuals per arm. We set the sample size liberally at 50 to allow for the possibility that we might be able to combine studies quantitatively in a meta-analysis.

## **Geographic Limit and Publication Dates**

Because health care systems, disparity indicators, and groups subject to disparities vary geographically, we limited eligible studies to those of patients in the U.S. health care system. Consistent with this inclusion criterion, only studies published in English were included. Searches were limited to papers published in 1983 or later because seminal work regarding QI strategies began to be published in the early 1980s.

# **Study Design**

We included randomized controlled trials (RCT), including cluster randomized controlled trials; controlled trials, including quasi-randomized studies; controlled before-after studies; prospective and retrospective cohort studies; interrupted time series with comparison groups; and stepped wedge designs.

## **Study Groups**

All studies were required to include a comparison group that did not receive the QI intervention or that received a different intervention. In addition, they were required to provide data that could be used to measure a disparity based on one of the population characteristics specified in the protocol before and after the intervention. These data could have included reference to an external reference group, but if they used an external referent group, the data needed to have been collected within four years of the enrollment of the target group and be from a source that was at the state or local level.

### **Interventions**

We considered both formal QI models and QI strategies for the review. As a starting point, we borrowed the taxonomy of QI strategies described in the antecedent Closing the Quality Gap series. To this taxonomy, we added a generic category (i.e., "other") to ensure retrieval of studies describing a QI strategy not specifically captured by the taxonomy. We did not include papers describing topics or interventions covered by other reports in the Closing the Quality Gap series (e.g., studies targeting public reporting, payment bundling, and medication adherence).

### **Conditions**

We sought studies of interventions to reduce disparities in health and process outcomes associated with a targeted set of clinical conditions, namely:

- Asthma
- Cancer:
  - Colorectal cancer (including screening)

- o Breast cancer (including screening)
- Cardiovascular disease:
  - o Congestive heart failure
  - o Coronary artery disease (including ischemic heart disease, myocardial infarction, and acute coronary syndrome)
  - o Hypertension
- Cystic fibrosis
- Depression (major depressive disorder only)
- Diabetes
- End stage renal disease
- Pneumonia (including pneumococcal vaccination)
- Pregnancy

The selection of these conditions was based on priority lists previously published by AHRQ and the Institute of Medicine (IOM)<sup>12</sup> and through consultation with the TEP.

### **Outcomes**

We included studies that captured health outcome measures and/or process measures to answer Key Question (KQ) 1. For KQ 2, we sought studies that reported harms (e.g., negative unintended consequences, misallocation of effort, decreased patient satisfaction, etc.) of the QI intervention to individual participants or the health care system. Table 2 summarizes the inclusion and exclusion criteria.

Table 2. Inclusion and exclusion criteria

Category	Criteria	
Population	Individuals receiving health care in the United States for a pre-specified clinical condition:	
	Studies had to include data on these characteristics of the study population known to be associated with health disparities: race or ethnicity, socioeconomic status, insurance status, sex, sexual orientation, health literacy/numeracy, and/or language barrier.	
Intervention	QI strategy: (1) a formal, broad organizational model; or (2) a change process in health care systems, services, or supplier for the purpose of increasing the likelihood of optimal clinical quality of care.	
Comparator	Usual care or use of an alternate strategy.	
Outcome(s)	Outcome measures of interest included: health outcome measures (e.g., morbidity and mortality, indirect health outcomes such as blood pressure and HbA1c); process measures (e.g., proportion of patients treated according to clinical guidelines); changes in disparity; and harms (i.e., any negative impact of the intervention to the individual patients or the health care system).	
Time period	1983-present	
Setting	Strategies conducted in or based out of a hospital, provider office, and/or health care clinic.	
Other criteria	<ul> <li>Admissible designs included: randomized controlled trials, including cluster randomized controlled trials; controlled trials, including quasi-randomized trials; controlled before-after studies; prospective and retrospective cohort studies; interrupted time series studies with comparison groups; and stepped-wedge design studies.</li> <li>Original research studies with sufficient detail to enable use and adjustment of the data and results.</li> <li>Inclusion of a target group and an internal or external referent group to measure changes in disparities.</li> <li>A minimum sample size of 50 individuals per study and intervention group or subgroup.</li> <li>Extractable data on relevant outcomes from text or tables.</li> <li>English language publications only</li> </ul>	

# **Screening of Studies**

We conducted screening in two phases: abstract and full text screening. Two reviewers independently reviewed each abstract. All papers with inclusion—exclusion conflicts at the abstract review level or lacking adequate information to make a determination were promoted to full text review. Two reviewers independently reviewed the full text of papers included at the abstract phase. Disagreements between reviewers at the full-text screening level were resolved by a senior investigator.

## **Data Extraction and Management**

#### **Evidence Tables**

Two reviewers independently extracted relevant data (e.g., setting, condition, patient population, QI strategy, outcomes, and disparity) from all included papers using a predefined evidence table shell. A senior investigator reviewed the evidence table for accuracy and completeness. The research team met regularly during the data extraction period and discussed global issues related to the process. The final evidence table is presented in Appendix I. When possible to identify, analyses resulting from the same study were grouped together.

### **Statistical Tests**

When provided in the study paper, we report p values for tests of significance and confidence coefficients with confidence intervals for reliability of estimates in the evidence tables and results section of the report. In rare cases, the report authors calculated the statistical significance of study results using a standard statistical test (Fisher's Exact Two-Tailed Test). These are indicated in footnotes.

## **Quality Assessment of Studies**

To interpret study results and grade the strength of evidence, we assessed the methodological quality of individual studies using two existing tools. Senior investigators assigned a study design to each of the included papers according to the description of the study methods. Two investigators independently assessed the quality of individual studies. A senior investigator resolved discrepancies between reviewers.

### **Quality Assessment Tools**

We assessed quality for each outcome of interest from included studies using design-specific criteria for RCTs and cohort studies. To assess internal validity of RCTs, we used the Cochrane Collaboration Risk of Bias tool, which evaluates domains including sequence generation, allocation concealment, blinding, outcome data reporting, and reporting bias (see Appendix D and Appendix E).

For nonrandomized and observational studies, we used the Newcastle-Ottawa scale<sup>13</sup> to assess three broad perspectives: (1) the selection of the study groups; (2) the comparability of the groups; and (3) the ascertainment of either the exposure or outcome of interest for case-control or cohort studies, respectively (see Appendix F).

The Newcastle-Ottawa Quality Assessment Scale includes eight multiple choice questions from three broad domains: four items related to selection of cohorts, one item related to comparability of cohorts, and three items related to assessment of outcomes. <sup>13</sup>

# **Determining Quality Ratings**

These risk of bias scoring and quality assessment tool ratings were based upon study design and conduct (i.e., internal study validity). We rated individual studies as good, fair, or poor and included data from good and fair quality studies in our analysis. We designated thresholds to rate individual studies as "good," "fair," or "poor" quality. Appendix G outlines requirements for each rating.

Several of the included papers reported data from a post hoc or secondary analysis of a previously completed RCT. Because the balance between groups achieved by randomization does not reliably extend to subgroups, the team modified the risk of bias/quality assessment on a case-by-case basis, considering the methods of the individual paper and parent study methods when appropriate.

## **Data Synthesis**

Meta-analysis was not appropriate in this review due to the substantial heterogeneity of study interventions, patient populations and outcomes; therefore, all analysis is narrative and based on evidence and summary tables. Studies are summarized by the disparity addressed in the research and in categories of clinical conditions. We use the race and ethnicity categorizations that were used by the authors of the included studies.

## Grading the Strength of Evidence for a Body of Evidence

The assessment of the literature is done by considering both the observed effectiveness of interventions and the confidence that we have in the stability of those effects in the face of future research. The degree of confidence that the observed effect of an intervention is unlikely to change is presented as strength of evidence, and it can be regarded as insufficient, low, moderate, or high. Thus, strength of evidence describes the adequacy of the current research, both in terms of quantity and quality, as well as the degree to which the entire body of current research provides a consistent and precise estimate of effect. Interventions that have demonstrated benefit in a small number of studies but have not yet been replicated using the most rigorous study designs will therefore have insufficient or low strength of evidence to describe the body of research. Future research may find that the intervention is either effective or ineffective. Interventions that demonstrate consistent results from well-designed studies that minimize risk of bias will be assessed as high or moderate depending upon the likelihood that future research will alter the current evidence.

Methods for applying strength of evidence assessments are established in the EPC's Methods Guide for Effectiveness and Comparative Effectiveness Reviews<sup>14</sup> and are based on consideration of four domains: risk of bias (low, medium, high), consistency (inconsistency not present, inconsistency present, unknown or not applicable), directness (direct, indirect), and precision (precise, imprecise). We assessed strength of evidence separately for each major intervention-outcome pair. We assigned an overall evidence grade to each key outcome for each comparison of interest based on the ratings for the individual four domains. Once we had established the maximum strength of evidence possible based upon these four domains we assessed the number of studies and range of study designs for a given intervention-outcome pair, and downgraded the rating when the cumulative evidence was not sufficient to justify the higher rating. The possible grades were:

- High: High confidence that the evidence reflects the true effect. Further research is unlikely to change estimates.
- Moderate: Moderate confidence that the evidence reflects the true effect. Further research may change our confidence in the estimate of effect and may change the estimate.
- Low: Low confidence that the evidence reflects the true effect. Further research is likely to change confidence in the estimate of effect and is also likely to change the estimate.
- Insufficient: Evidence is either unavailable or does not permit a conclusion.

# **Peer Review and Public Commentary**

Experts in the fields of QI and health disparities and individuals representing stakeholder and user communities were invited to provide external peer review. AHRQ and an associate editor also provided comments. The draft report was posted on the AHRQ Web site for 4 weeks to elicit public comment. We addressed all reviewer comments and revised the report as appropriate. A disposition of comments report will be available 3 months after the Agency posts the final report on the AHRQ website.

### Results

This chapter presents the results of the review of quality improvement (QI) strategies to address health care disparities. We present findings for Key Question 1 (KQ 1) beginning with an overview of the content of the literature as a whole, followed by results and detailed analysis organized first by the disparity addressed in the research and then again by clinical conditions. No studies provided information on Key Question 2 (KQ 2), which pertained to harms associated with the interventions.

Studies also are described in more detail in summary tables in the relevant section of text. For information on quality scores for each study, see Appendix H; for information on the strength of evidence for outcomes, see Appendix J.

### Search Results and Included Studies

Using a broad search strategy, we identified 4,278 titles and abstracts with potential relevance for initial screening (Figure 2). From this screening, 791 papers were identified as possibly related and moved forward for full-text review. At this second level, 772 papers did not meet eligibility criteria. The complete list of exclusion reasons is provided in Appendix L.

Therefore, for this report, we synthesized data from 19 papers representing 14 unique studies that addressed the following conditions: breast and colorectal cancer screening (n=5); cardiovascular disease, including coronary heart disease and hypertension (n=5); depression (n=3); and diabetes (n=3).

All papers addressed KQ 1. The included studies used an internal referent group to measure disparity; none of the included papers relied upon an external referent group to measure disparities.

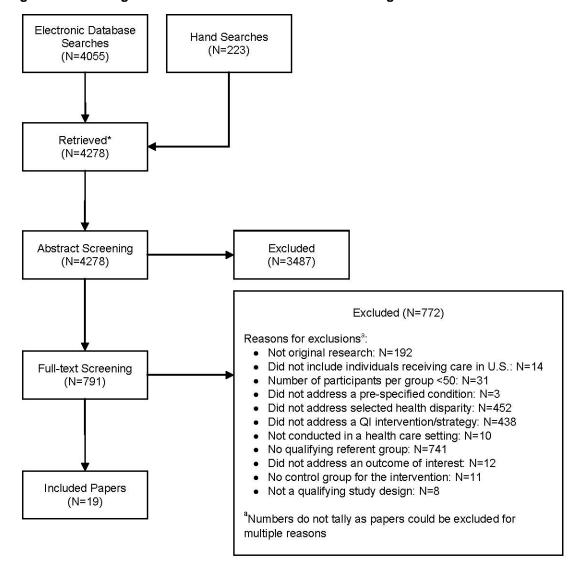


Figure 2. Flow diagram of the literature search and screening

 $\textbf{Abbreviation:} \ QI = quality \ improvement.$ 

## **Overview of Included Studies**

Nineteen papers qualified for our review (Table 3), reflecting results from 14 unique studies. Results are further summarized with details in the evidence tables (Appendix I).

Of the 14 studies, 11 were randomized controlled trials (RCTs), <sup>15-26</sup> including 2 cluster RCTs. <sup>15,16,27-30</sup> The remaining studies were cohort studies, including one prospective cohort study, <sup>31</sup> one retrospective cohort study, <sup>32</sup> and one cohort study with a historical control. <sup>33</sup>

Included papers targeted or described disparities associated with differences in race or ethnicity (n=14), <sup>15-17,21,22,24,28,29,31-34</sup> socioeconomic status (n=3), <sup>18,24,32</sup> insurance status (n=2), <sup>19,26</sup> language (n=2), <sup>20,26</sup> health literacy (n=1), <sup>23</sup> and sex (n=1). All studies were focused in a specific clinical area. Five studied a QI intervention in cancer screening, three in cardiovascular disease, three in depression care, and three in diabetes.

Outcomes included health care processes and health outcomes. All but one of the studies incorporated multiple components into their QI approach. Patient education was a part of most

interventions (12 of 14), although the specific approach differed substantially across the studies. Ten of the studies incorporated self-management; this would, include, for example, teaching individuals with diabetes to check their blood sugar regularly. Most (8 of 14) included some sort of provider education, which may have focused on the clinical issue or on raising awareness about disparities affecting the target population. Nonetheless, given the degree to which the interventions all included multiple components that were implemented as a system, it is not possible to tease apart the effects or implications of individual aspects.

We organize the results in two ways. First, the results are summarized by effect on particular disparities, which is the primary focus of the review; second, descriptions of the studies are organized by clinical condition as a reference for end users interested in implementing QI approaches in individual clinics or clinical specialties.

Table 3. Overview of the studies of QI interventions to address disparities in health outcomes

Characteristic	Race or Ethnicity	Insurance Status	Language Barrier	Health Literacy	Socioeconomic Status	Sex
Intervention category						
Patient Education	10	1	2	1	2	1
Provider Education	6	0	0	1	2	1
Promotion of Self-management	10	1	2	0	2	1
Audit and Feedback	3	0	0	1	0	1
Facilitated Relay of Clinical Data to Providers	2	0	0	0	1	1
Patient Reminder System	2	2	2	0	0	0
Provider Reminder System	1	0	0	0	1	0
Organizational Change	1	0	0	0	0	0
Other	5	0	0	0	2	1
Condition						
Cancer	2	2	2	1	0	0
Cardiovascular Disease	2	0	0	0	0	0
Depression	3	0	0	0	2	1
Diabetes	3	0	0	0	0	0
Design						
RCT	9	2	2	1	2	1
Retrospective cohort	2	0	0	0	0	0
Prospective cohort	1	0	0	0	0	0

**Abbreviation:** RCT = randomized controlled trial.

# **Effects on Racial or Ethnic Disparities**

#### Overview

Eleven studies provided data on the effects of QI interventions on disparities in health care associated with race or ethnicity (Table 4).

Two RCTs evaluated the effects of interventions on racial or ethnic disparities in colorectal cancer screening. <sup>25,26</sup> Three studies explored the effects of various QI strategies on disparities associated with race in cardiovascular disease, including coronary artery disease and hypertension. <sup>21,22,33</sup> Three studies evaluated the effect of QI interventions on racial or ethnic disparities in depression outcomes <sup>15,17,24,28,29,34</sup> and three studies provided data to assess the impact of QI interventions on racial or ethnic disparities in diabetes outcomes. <sup>16,31,32</sup>

Most (n=8) were unable to show a reduction in disparity, or suggest that they might be useful for reducing disparity by failing to demonstrate an amplified effect in any one subgroup. Three studies provided data suggesting that their approach had some potential to affect disparities. None provided conclusive information and each studied a different approach in a different disease condition, making synthesis challenging.

One disease management and patient education program<sup>32</sup> was associated with a reduction in disparity between Black and White patients in HbA1c testing when it was targeted in a geographic area with very high rates of diabetes. Other interventions did not demonstrate a significant reduction in disparity, but an amplified effect was seen in nonwhite populations with some interventions, including a patient education program to reduce blood pressure, <sup>21</sup> and a complex collaborative care model targeted to providers of patients with depression. <sup>15,28,29,34</sup> In the latter study, the intervention was significantly more effective in the short term among minorities, although this was no longer the case at 1 year, and the intervention was not effective in any population at five and nine years.

Table 4. Summary of effects on disparities in health outcomes associated with race or ethnicity

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Arean et al., 2005 <sup>17</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>No disparity in depression severity existed by race or income at baseline.</li> <li>The intervention was effective in all racial subgroups with no interaction by race and no amplified effect in any group.</li> <li>In subgroup analysis, the intervention was associated with greater use of psychotherapy but not pharmacotherapy within the Black population</li> </ul>
Bao et al., 2011 <sup>24</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Provider reminder system</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>At baseline within the usual care group, 22% of minorities had adequate antidepressant use, compared to 39% of White patients.</li> <li>The intervention had no effect on this disparity and ethnic minorities did not receive greater benefit from intervention compared with White patients during any time period.</li> </ul>

Table 4. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Bosworth et al., 2011 <sup>21</sup> RCT Cardiovascular disease: hypertension	<ul> <li>Patient education</li> <li>Promotion of self- management</li> </ul>	<ul> <li>The race by time by treatment group effect model suggested differential intervention effects on blood pressure (BP) over time for White patients vs. nonwhite patients for both systolic blood pressure (SBP) (p=0.08) and diastolic blood pressure (DBP) (p=0.01).</li> <li>The combination of home BP monitoring and tailored behavioral intervention was most effective in nonwhite participants at 24 months (p=0.04).</li> </ul>
Coberley et al., 2007 <sup>32</sup> Retrospective cohort Diabetes	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Organizational change (disease management)</li> </ul>	<ul> <li>Initial racial disparity in HbA1c testing between the health disparity zone (HDZ) group and non-HDZ group was 12%.</li> <li>Disparity was not significantly reduced after 12 months (p=0.06).</li> <li>Within the HDZ zone (high prevalence of diabetes), testing increased by15% among Black participants but not among White participants, resulting in a reduction in disparity in this subgroup analysis.</li> </ul>
Connett and Stamler, 1984 <sup>22</sup> RCT Cardiovascular disease: coronary artery disease and hypertension	<ul> <li>Patient education</li> <li>Promotion of self-management</li> </ul>	<ul> <li>At baseline, Black participants had higher rates of smoking than White participants (68.7% vs. 63%; p&lt;.001).</li> <li>Both racial groups experienced significant reductions in smoking of close to 50% in the intervention group and more than 35% in the usual care group.</li> <li>The baseline disparity persisted in the intervention group but was apparently reduced in the usual care group.</li> <li>A statistically significant but clinically insignificant disparity in DBP and SBP by race was present at baseline.</li> <li>Blood pressures were reduced in both the intervention and control groups, with greater change observed in the intervention group.</li> <li>The small disparity observed at baseline was further reduced at followup in the intervention group but not the control group.</li> </ul>
Lasser et al., 2011 <sup>26</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates at baseline by race or ethnicity.</li> <li>The intervention was more effective in White and Black individuals relative to those of other or unknown race.</li> </ul>

Table 4. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Mahotiere et al., 2006 <sup>31</sup> Prospective cohort Diabetes	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Other (community intervention)</li> </ul>	<ul> <li>The disparity in biennial lipid profile testing at baseline was 19%.</li> <li>The biennial lipid profile testing rate improved by 26.2 percent in African-American beneficiaries with diabetes in the intervention areas following implementation of the QI program.</li> <li>The disparity in performance of biennial lipid profile between African-American Medicare fee-for-service beneficiaries and White Medicare fee-for-service beneficiaries reduced to 9.2% following implementation of the QI program.</li> <li>An analysis of the direct impact of the selected interventions on reducing the disparity in this uncontrolled database analysis was not feasible.</li> </ul>
Miranda et al., 2003 <sup>15</sup> ; Miranda et al., 2004 <sup>34</sup> ; Wells et al., 2007 <sup>28</sup> ; Wells et al., 2004 <sup>29</sup> Cluster RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>The intervention was associated with decreases in probable depressive disorder among minorities but not White patients at 12 months. (Latino p=0.02; African-American p=0.01).</li> <li>At 12 months among intervention recipients, the baseline disparity increased (6.7% to 7.7%) between Latino and White patients, and decreased between African-American and White patients (9.2% to 6.7%).</li> <li>Although a statistically significant interaction was seen between intervention and ethnicity at 6 months when minorities were grouped and contrasted with White patients, no such interaction persisted at 12 months.</li> <li>The overall effect of the intervention on depression status was not significant at five and nine years, but an interaction with race was seen in the overall model of effectiveness. The intervention was associated with improvements in the Mental Health Inventory among minorities (p=0.008) but not among White patients (p=0.59).</li> <li>In subanalysis at five years, QI-Therapy but not QI-Meds was effective within the minority population.</li> </ul>
Olomu et al., 2010 <sup>33</sup> Retrospective cohort (historic controls) Cardiovascular disease: coronary artery disease	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (guideline adherence)</li> </ul>	<ul> <li>The American College of Cardiology's Acute Myocardial Infarction (AMI) Guidelines Applied in Practice (GAP) strategy was associated with increased inpatient use of beta-blockers among nonwhite patients.</li> <li>Racial disparities in use of cardiac catheterization and percutaneous coronary intervention appeared after implementation of the GAP QI strategy despite overall improvements in care.</li> <li>The admission tool and inpatient aspirin were more often used post-GAP than pre-GAP in both White and nonwhite patients.</li> </ul>
Sequist et al., 2010 <sup>16</sup> Cluster RCT Diabetes	<ul><li>Provider education</li><li>Audit and feedback</li></ul>	<ul> <li>Disparities were present at baseline in HbA1c levels, BP control, and LDL level.</li> <li>The intervention showed no effect overall in either racial group.</li> <li>The intervention did not reduce the disparity.</li> </ul>

Table 4. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Siddiqui et al., 2011 <sup>25</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates at baseline by race or ethnicity.</li> <li>No statistically significant difference in screening rates between White participants and African-Americans in the control group.</li> <li>When intervention groups were combined, the screening rate was significantly higher in White participants compared to African-Americans.</li> </ul>

**Abbreviations:** BP = blood pressure; CRC = colorectal cancer; DBP = diastolic blood pressure; GAP = American College of Cardiology's AMI Guidelines Applied in Practice; HbA1c = hemoglobin A1c; HDZ = health disparity zone; LDL = low density lipoprotein; QI = quality improvement; RCT = randomized controlled trial; SBP = systolic blood pressure.

### **Detailed Description**

### **Studies Focused on Cancer Outcomes**

Two studies evaluated the effects of interventions on racial or ethnic disparities in colorectal cancer screening. <sup>25,26</sup> Both studies were secondary analyses of RCTs and examined the usefulness of patient education and reminder systems for improving colorectal cancer screening rates, as measured by participation in fecal occult blood testing or colonoscopy within one year after study entry. One study involved a primary care practice-based research network <sup>26</sup> and the other included a single large academic primary care practice. <sup>25</sup> Both trials took place in the United States and targeted patients who were due for CRC screening. The two studies employed a usual care comparison group.

One of the secondary RCT analyses randomized 465 participants, including 221 White patients, 129 Black patients, and 115 patients of other/unknown race. <sup>25</sup> Patients were assigned to either usual care or language-concordant assistance by a patient navigator, a trained individual who promoted self-management strategies and provided patient education and reminders. The patient navigator intervention as compared with usual care was associated with significantly increased screening rates among both White (33.9 vs. 16.5 percent, p=0.003) and Black participants (39.7 vs. 16.7%, p=0.004), with overall similar proportions obtaining screening in both racial groups during the 12 month followup period.

The other secondary RCT analysis randomized 1430 patients, including 578 Whites and 852 Black participants. <sup>26</sup> This trial included a usual care comparison group and employed three intervention arms: a standard intervention including an educational brochure and screening reminder, a tailored intervention including the standard materials plus additional educational information targeting each participant's perceived barriers to screening as assessed at baseline, and a group that received both the tailored intervention plus a reminder phone call. CRC screening rates by 12-month followup were similar in the usual care group for Whites as compared with African-Americans (33 percent vs. 32 percent). An analysis pooling results of the three intervention groups vs. usual care indicated that the interventions had an increased effect on improved screening rates among White participants as compared with their Black counterparts (53 percent vs. 43 percent, adjusted OR 1.44, 95% CI 1-12 to 1.86, p=0.005).

### **Studies Focused on Cardiovascular Disease**

### **Coronary Artery Disease**

One post hoc analysis of an RCT,<sup>22</sup> and one retrospective cohort study<sup>33</sup> explored the effects of various QI strategies on racial health care disparities in coronary artery disease. The RCT addressed reduction of associated risk factors,<sup>22</sup> while the retrospective cohort examined management of AMI.<sup>33</sup> QI strategies included patient education and facilitation of self–management<sup>22</sup> and a multifactorial provider– and systems–focused strategy.<sup>33</sup> Both studies took place in the United States, in the setting of two multicenter collaborations of academic and community health centers.<sup>22,33</sup> The studies each employed an internal usual care comparison group.

For coronary artery disease (CAD), we sought assessments of critically important clinical outcomes, surrogate clinical outcomes, and process measures. No health outcomes (e.g., death, myocardial infarction, myocardial ischemia, and congestive heart failure were reported. One surrogate outcome in the form of a clinical risk factor for CAD (smoking) was reported. Several process measures (cardiac catheterization, PCI, beta-blocker usage, aspirin usage) were reported.

One analysis of RCT data<sup>22</sup> evaluated the utility of QI strategies in reducing CAD risk factors and one retrospective cohort study<sup>33</sup> evaluated QI strategies for improving management of acute myocardial function among different racial groups.

The RCT was the Multiple Risk Factor Intervention Trial (MRFIT), a large secondary prevention study among men at elevated risk of CAD. The intervention included an initial intensive period of ten weekly sessions followed by individual counseling, focused on behavior change and risk factor reduction related to nutrition, smoking cessation, and antihypertensive therapies. The study recruited men from 22 academic and community health care centers in the United States. A secondary analysis examined racial differences in CAD risk factor reduction in 11935 White patients (including individuals who identified their race as "other") and 931 Black patients. The sixth annual followup visit included 5754 in the intervention arm (5338 White patients and 416 Black patients) and 5638 in the usual care group (5227 White patients and 411 Black patients). The incidence of smoking cessation (self–report adjusted for thiocyanate levels) by the sixth annual visit was approximately the same among White and Black patients in the intervention group, 46.0 percent and 43.0 percent, respectively (p=0.26). In the usual care group at this same time point, Black patients were less likely to have quit smoking than White patients, at 22.5 percent versus 29.0 percent (p<0.004).

One fair retrospective cohort study, using historical controls<sup>33</sup> constructed a retrospective cohort from Medicare patients with history of AMI treated at a group of academic and community hospitals in Michigan, sampling data from one year preceding a rapid cycle QI project and from four months immediately following implementation. The QI strategy involved customization and implementation of the American College of Cardiology's AMI Guidelines Applied in Practice (GAP) toolkit,<sup>36</sup> including provider education, standing orders, admission and discharge tools, and pocket guidelines for clinicians. The analysis compared results between the group of patients (including 1158 White and 210 nonwhite patients) treated before implementation of the GAP toolkit (pre-GAP), and a separate group of patients (including 1209 White and 280 nonwhite patients) treated after implementation of the GAP toolkit (post–GAP). A significant improvement in use of the admission tool and aspirin during the inpatient stay were observed in both White and nonwhite patients. Nonwhite patients had a significant increase in

<sup>\*</sup>Fisher's Exact Two-Tailed Test, calculated by review authors using original data.

inpatient beta-blockade (66.0 percent pre-GAP vs. 83.3 percent post-GAP, p=0.04), while White patients did not (76.8 percent pre-GAP vs. 82.4 percent post-GAP, p=0.13).

Although outcomes improved in other measures for both racial groups, they improved more for White patients, creating or worsening the disparity between the two groups, while improving outcomes overall. For example, pre-GAP, use of the discharge tool was low overall, but similar for White and nonwhite patients (1.8 percent vs. 1.0 percent, p=0.37). After implementation of GAP, the tool was used more overall, but significantly more often with White patients than nonwhite patients (30.1 percent vs. 23.6 percent, p=0.03). A similar effect was observed for smoking cessation counseling, which was similar pre-GAP among White and nonwhite patients (39.4 percent vs. 34.2 percent, p=0.57), but post-GAP, increased significantly more among White patients (73.4 percent vs. 50.0 percent, p=0.002).

Disparities also developed in use of invasive procedures. For example, the use of cardiac catheterization, which was similar among White and nonwhite patients pre-GAP, increased significantly among White patients post-GAP (from 45.5 percent to 50.8 percent, p=0.01), though it tended to decrease among nonwhite patients (from 44.8 percent pre-GAP to 36.8 percent post-GAP, p=0.07). Among White patients, PCI increased from 19.8 percent to 25.6 percent, p=0.0008; however among nonwhite patients, PCI remained flat (15.7 percent pre-GAP vs. 13.2 percent post-GAP, p=0.44). Consequently, in the post-GAP period, use of these invasive procedures was significantly less common among nonwhite patients than White patients (for cardiac catheterization, OR 0.56, 95% CI, 0.43 to 0.74; for PCI, OR 0.44, 95% CI, 0.31 to 0.64 In summary, this study of the GAP initiative demonstrated mixed results regarding disparities in cardiovascular care. Use of the admission tool and inpatient aspirin increased similarly among White and nonwhite patients; use of inpatient beta-blockade increased more among nonwhite patients; and use of cardiac catheterization and PCI increased more among White patients than nonwhite patients.

### Hypertension

Two post hoc analyses of RCTs explored the effects of various QI strategies on racial disparities in hypertension. <sup>21,22</sup> The RCTs addressed management of hypertension and reduction of risk factors for CAD including hypertension. <sup>22</sup> QI strategies were patient education and facilitation of self–management. <sup>21,22</sup> Both studies took place in the United States, with settings including university clinics <sup>21</sup> and multicenter collaborations of academic and community health centers. <sup>22</sup> The studies each employed an internal usual care comparison group.

Health outcomes such as death, myocardial infarction, ischemic stroke, and congestive heart failure were not reported. One surrogate clinical outcome (blood pressure) was reported.

Two post hoc analyses of RCT data assessed the impact of QI strategies on hypertension among different racial groups. The MRFIT trial, described above for its modification of coronary artery disease risk factors, assessed the effect of the intervention on systolic and diastolic blood pressure among Black and White participants. He neceived monitoring of blood pressure and lifestyle counseling; upon being diagnosed with hypertension, they received stepped pharmacotherapy. The study quality was assessed as fair; the data presented here are from post hoc analyses.

At baseline, Black men had higher mean blood pressure than White men (138/94 vs. 135/91, p<0.001). The intervention was reported to have resulted in slightly greater reduction of blood pressure among Black men (reduced systolic blood pressure [SBP] by 11.7 percent and diastolic blood pressure [DBP] by 13.6 percent), compared with White men (reduced SBP by 10.3 percent

and DBP by 11.4 percent).\*\* In the control group, less reduction was observed compared with the intervention, and similar changes in blood pressure were reported among Black men (reduced SBP by 5.8 percent and DBP by 8.1 percent) and White men (reduced SBP by 6.4 percent and DBP by 7.9 percent).\*\* At the sixth annual followup visit, among intervention patients, mean blood pressures were reported to be similar in Black (122/81) and White (121/80) men, and a similar proportion of Black (72 percent) and White (71 percent) participants were at or below the DBP goal. Achievement of these goals was not related to baseline DBP. Among control group patients, mean blood pressure was reported as slightly higher among Black men (129/86), than White men (127/84).\*\* Blood pressures in the intervention group were reduced more than were those in the control group. A small, but significant baseline disparity in systolic and diastolic blood pressure measures was not present at followup in the intervention group, but persisted at followup in the control group.

Another post hoc analysis<sup>21</sup> explored racial differences in outcomes following an RCT that assessed two hypertension self–management strategies (home blood pressure [BP] monitoring 3 times weekly, or tailored behavioral self–management intervention by phone every other month) alone or in combination, compared with usual care. Participants included White (n=308) and nonwhite (n=328, 95 percent African-American) patients with a diagnosis of hypertension for at least 1 year before recruitment.<sup>21</sup> Participants were treated in two university–affiliated general internal medicine clinics and followed for 24 months. The study quality was assessed as fair; the data presented here are from a post hoc analysis. At baseline, nonwhite patients had significantly higher SBP (128 vs. 121 mm Hg, p<0.0001) and DBP (74 vs. 69 mm Hg, p<0.0001) than White participants, and the initial prevalence of blood pressure control was also lower among nonwhite than White patients (72.5 vs. 88 percent, p<0.0001). A race by time by treatment group effect model suggested differential intervention effects on BP over time for White and nonwhite patients for DBP (p=0.01), but not for SBP (p=0.08).

Among White patients, SBP was not significantly different at 12 or 24 months followup in intervention and control groups. Results for DBP were similar, except mean DBP at 1 year was 3 mm Hg higher in the behavioral only strategy, compared with control (p=0.03). Among nonwhite patients, at 1 year, SBP and DBP improved by 5 mm Hg and 3 mm Hg, respectively, in each of the three intervention groups, compared with usual care (p<0.05). In the combined intervention group (home monitoring + tailored behavioral intervention), persistent improvements in SBP (-7.5 mm Hg) and DBP (-3.5 mm Hg) were present at 24 months, compared with usual care (p<0.05). Thus, the two hypertension self-management strategies were ineffective among White patients, but effective among nonwhite patients. The interventions alone or in combination significantly reduced BP at 1 year in the nonwhite group, and the combined intervention resulted in persistent BP reductions at 24 months, thereby reducing disparities in BP control.

### **Studies Focused on Depression Outcomes**

Three studies evaluated the effect of QI interventions on racial or ethnic disparities in depression outcomes. 15,17,24,28,29,34 All three studies used a collaborative care model, which involved collaboration among multiple clinical providers to provide a coordinated set of interventions. The clinical model in all three studies included a dedicated mental health coordinator (nurse or case manager), creation of mental health teams (composed of primary provider, facility nurses, and psychiatrists), evidence-based pharmacotherapy and psychotherapy, extensive provider education, and longitudinal patient followup to evaluate clinical status and

<sup>\*\*</sup>The authors did not provide statistical tests of significance, and the standard deviations for the means were not reported, so statistical analysis could not be performed by the reviewers.

adherence. Each intervention was designed to address known barriers to the receipt of quality mental health care.

All three trials were prospective, randomized controlled trials, with randomization to types of provider training occurring at the practice level. However, given the nature of the intervention (group-level randomization), individual providers and patients retained ability to select the treatment provided to the individual patient. All three trials took place in the United States.

The Partners in Care (PIC) study <sup>18,21,31,34,35,41</sup> delivered a composite intervention of patient

The Partners in Care (PIC) study<sup>18,21,31,34,35,41</sup> delivered a composite intervention of patient and provider education, nurse-assisted patient assessment, and targeted use of medication management and cognitive behavioral therapy. Outcomes were collected through mail surveys at 6 months, <sup>34</sup> 1 year, <sup>15</sup> 2 years, <sup>30</sup> 5 years<sup>29</sup> and 9 years. <sup>28</sup>

The study enrolled 1356 patients, and was conducted in six managed care organizations around the United States (at least one organization was selected in each of the four United States' census regions; study investigators also sought to recruit organizations known to have high enrollment of Mexican Americans).<sup>37</sup>

Study patients were selected by use of the 12-month Composite International Diagnostic Interview, version 2.1, an unspecified tool designed to evaluate depressed symptoms in the prior month, telephone interviews, and self-administered questionnaire. Fourteen percent of the patients screened for the study were eligible to participate. Of those eligible, 34.7 percent enrolled. The study population included 61.3 percent Caucasians, 30 percent Latinos, and 7.3 percent African-Americans.

PIC was designed to compare usual care versus two interventions: QI–Meds and QI–Therapy. The intervention components: (1) enrolled practices provided in-kind resources (up to one half of the cost) to assist in intervention implementation; (2) provider training at study onset and monthly meeting thereafter to provide feedback on treatment patterns; (3) each practice had a designated staff nurse who was trained as a "depression specialist" who would screen patients for depression, then educate and engage those who screened positive in depression selfmanagement. The QI–Meds intervention contained all above components plus trained nurses who assessed patient adherence to antidepressant pharmacotherapy. The QI–Therapy intervention contained all of the above components plus local psychotherapists who could provide individual or group cognitive behavioral therapy.

The QI educational material was provided to patients in English or Spanish language text. Minority primary care providers were featured in videotapes viewed by intervention participants. Providers in intervention practices were given instruction in how to lower barriers to treatment experienced by Latino and African-American patients. PIC investigators, belonging to ethnic minority subgroups, also provided direct supervision for local experts.

At 1 year,<sup>15</sup> the ethnic minorities in the intervention group (QI–Meds and QI–Therapy were combined) had statistically significantly lower rates of probable depression than minorities receiving usual care (Latino 27 percent less, p=0.02; African-American 27 percent less, p=0.01). Conversely, no such effect was observed among White participants. The disparity between Latino participants and the White participants who received the intervention was reported to have increased in the first year of participant followup (6.7 percent at baseline to 7.7 percent at 1 year).\*\* In contrast, the disparity was reported to have decreased when comparing African-Americans to White participants (9.2 percent to 6.7 percent).\*\* However, although a statistically significant interaction was seen between intervention and ethnicity at 6 months when minorities were grouped and contrasted with White participants, no such interaction persisted by 1 year.

<sup>\*\*</sup>The authors did not provide statistical tests of significance, and the standard deviations for the means were not reported, so statistical analysis could not be performed by the reviewers.

At five years, the PIC group provided aggregate and individual results for the two ethnic minorities (Latinos and African-Americans). In the aggregated analysis, minorities receiving QI–Therapy had a lower rate of probable depression compared with usual care minorities (35.6 percent vs. 55.8 percent, p=0.01). On the other hand, no statistically significant difference was observed in outcomes of QI–Meds in terms of probable depressive disorder in the minority population (45.4 percent vs. 55.8 percent, p=0.13).

The overall effect of the intervention on depression status was not significant at five and nine years. No disparity was established at baseline, and the data did not demonstrate a change in disparity associated with the intervention. <sup>28</sup>

Employment status did not change significantly among minorities as a result of the intervention compared with usual care employment rates (Latino 3.5 percent increase, p=0.38; African-American 5.3 percent decrease, p=0.43). A decrease in the disparity in employment between nonwhites and Whites decreased at 6 months from 10 percent to 3.2 percent, but the concordant gap between White participants and the Latino participants increased from 3.6 percent to 5.9 percent at 1 year of followup. The employment status of Black intervention recipients also worsened in the first year of followup: at baseline, 74.1 percent Black participants versus 64.5 percent White participants were employed; at 1 year, 70.3 percent versus 66 percent were employed. None of these changes could be attributed to the intervention, however.

Appropriate care was defined as identifying a patient's need for care and providing guideline concordant treatments. At 1 year, <sup>15</sup> minority intervention subgroups experienced improvement in the rate of appropriate care received compared with non-intervention minority groups (Latino 26.6 percent increase, p=0.03; African-American 26.3 percent increase, p=0.33). The disparity between White participants and minority participants was not decreased.

The five year evaluation reported likelihood of unmet need, rather than rates of appropriate depression care, among ethnic minorities. Ethnic minority participants in the QI–Meds group were 11.8 percent less likely to have unmet need compared with White participants in the QI–Meds group; ethnic minority participants in the QI–Therapy group were 5.3 percent less likely to have unmet need compared with White participants in the QI–Therapy group at 5 years. Because the baseline incidence of probable unmet need was not disaggregated by ethnicity, no comment can be made on whether the disparity gap was narrowed by either intervention.

The PROSPECT study <sup>24</sup> involved 20 primary care practices and 396 patients, 60 years of age or older, from the Northeastern United States. The intervention included a practice-based depression care manager who offered treatment recommendations to the primary care provider based on a prespecified algorithm. While the treatment algorithm focused on a particular pharmacotherapeutic regimen, the provider or patient could elect to modify the recommended course. If pharmacotherapy was declined, then psychotherapy was recommended. Care managers were also involved with monitoring patient treatment response and adherence. Providers in practices randomized to usual care received videotaped and printed information on depression in older patients; they also received written communication that alerted them if a patient fulfilled diagnostic criteria for depression.

The 2011 publication described a post hoc analysis of data that initially evaluated intervention efficacy among all older depressed primary care patients included in the study population. <sup>24</sup> In the original publication, <sup>38</sup> investigators reported that the intervention demonstrated efficacy in reduction of suicidal ideation and depression symptoms in the study population. Since that publication, the PROSPECT investigators conducted a subgroup analysis of 134 ethnic minority participants (34 percent) versus 262 non-Hispanic White participants (66

percent), and a separate subgroup analysis of 146 college-educated (31 percent) versus 323 non-college educated participants (69 percent).<sup>24</sup>

For appropriate use of antidepressants (i.e., adequate dose), a baseline disparity was observed in the usual care group only, in which 22 percent of minorities had adequate antidepressant use, compared with 39 percent of non-Hispanic White participants. The effect on difference in this measure by race was not statistically significant at any point in time.<sup>24</sup>

The IMPACT trial was initially designed to evaluate whether a collaborative care model would improve clinical and functional outcomes in primary care patients being treated for depression. Subsequent subgroup analyses <sup>17,18</sup> were performed to evaluate whether this intervention could decrease outcome disparities among older ethnic minorities <sup>17</sup> and among older adults of higher and lower incomes <sup>18</sup>. IMPACT was carried out across five states involving 18 primary care clinics and eight health care organizations. The intervention was a collaborative, stepped care model that consisted of a depression care specialist (DCS) who coordinated (1) primary care provider education about evidence-based treatment, (2) a depression care manager working with the patient and primary care provider to activate patients in the management of their depression, (3) ongoing mood and medication monitoring, (4) brief psychotherapy (known as "Problem Solving Treatment of Primary Care;" PST-PC), (5) a clinical information tracking system, and (6) ready access to a psychiatrist for additional consultation.

The original study population consisted of 1801 patients, above the age of 60 years, who met criteria for major depression, dysthymic disorder, or both. Though the sample was not stratified by ethnicity at enrollment, representation of ethnicities enrolled in the two treatment groups was fairly balanced (intervention contained 51 percent of the study's White patients, 51 percent of the study's Black patients, and only 41 percent of the study's Latino patients, p= 0.07). The collaborative care intervention itself did not incorporate accommodations for cultural differences. Participant data was collected at baseline, 3, 6, and 1 year. 40

Ethnicity was not reported by the investigators to have been associated with any statistically significant differences in study subjects' baseline use of mental health treatment services.

Target responses were a decrease of 50 percent or more in the Hopkins Symptom Checklist-20 (HSCL-20) score from baseline, and treatment remission (i.e., HSCL-20 score less than 0.5). In the overall study population, at 1 year, 45 percent of intervention patients had a 50 percent reduction in depressive symptoms compared with 19 percent of the control group patients (OR 3.45, 95% CI, 2.71 to 4.38). The intervention was equally effective in all groups without cultural adaptation.<sup>17</sup>

No statistically significant differences existed between ethnic subgroups at baseline in health related functional impairment, assessed with the Short Form Physical Functioning Scale. At 1 year, all ethnic groups had small, statistically significant improvements in functional impairment and there was no difference in effect by race or ethnicity. <sup>17</sup>

Service use, defined as use of antidepressants or psychotherapy, did not differ statistically by participant ethnicity at baseline. <sup>17</sup> At 1 year, collaborative care minorities had accessed significantly more guideline-concordant depression services than minorities in the usual care group (use of antidepressant: 64 percent, 95% CI, 55 to 72 percent vs. 45 percent, 95% CI, 36 to 55 percent, p=0.003; use of psychotherapy: 37 percent, 95% CI, 28 to 47 percent vs. 13 percent, 95% CI, 6.5 to 19 percent, p=0.002). The effect of the intervention was thus similar across ethnic and income groups. <sup>17</sup> Although no statistical interaction was observed overall, there were some notable differences in magnitude of effect by race or ethnicity. In particular, in stratified analysis,

collaborative care was associated with greater use of psychotherapy among Black patients than usual care, but not with greater use of antidepressants.<sup>17</sup>

#### **Studies Focused on Diabetes Outcomes**

Three good quality studies provided data to assess the impact of QI interventions on racial or ethnic disparities in diabetes outcomes. One was a cluster RCT, one was a prospective cohort, and one was a retrospective cohort study. One was a patient reminder system that was a part of a larger diabetes management program, one was a comparison of community provider interventions with statewide programming, and the third was a program to provide cultural competency training to clinicians.

No studies reported health outcomes, including death, hypoglycemic coma, adverse drug event, cardiovascular complications, retinopathy progression, nephropathy progression, neurologic complications, or hospitalization for a complication of diabetes. Studies reported surrogate clinical outcomes, clinical risk factors for diabetes comorbidities, and process measures. Clinical outcomes reported in this literature were surrogate measures: achievement of hemoglobin A1C (HbA1c) of less than 7 percent, achievement of target blood pressure (BP), and achievement of target low density lipoprotein (LDL) cholesterol level.

One study provided data on all three surrogate outcomes. <sup>16</sup> This good quality cluster RCT was conducted in eight ambulatory health centers in eastern Massachusetts between 2007 and 2008 and randomized 124 primary care clinicians (physicians, nurse practitioners, and physician assistants) caring for 2699 (36 percent) Black diabetic patients and 4858 (64 percent) White diabetic patients to intervention or control. <sup>16</sup> Intervention clinicians received cultural competency training and monthly race-stratified performance reports that highlighted racial differences in control of HbA1c and LDL cholesterol levels and BP. Care recommendations for Black diabetic patients along with race-stratified physician-level diabetes performance reports were provided on a monthly basis to clinicians in the intervention group.

White and Black patients differed significantly at baseline for the rate of HbA1c less than 7 percent, with White patients more likely than Black patients to be controlled (46.1 percent vs. 40.0 percent, p<0.001) despite no disparity in receipt of annual HbA1c examinations. At followup, no intervention effect was observed in the Black patient population, with the proportion meeting the target in the intervention group just 3.3 percent higher than in the control group (adjusted difference 3.3 percent, 95% CI,-2.1 to 8.6, p= 0.24). By the same token, no intervention effect was seen in the White participants (adjusted difference 1.9 percent, 95% CI, -1.9 to 5.6, p= 0.22), and no effect was observed on the disparity between the two groups.

The same pattern was repeated for blood pressure and LDL outcomes, in which Black participants had consistently poorer health indicators, and the intervention was not shown to be effective in the Black population, and had no impact on disparities.

Process measures assessed in the diabetes QI literature were completion of HbA1c testing and lipid testing. A good quality retrospective cohort used a corporate national health care database to assess the effectiveness of disease management programs on testing for HbA1c levels in socioeconomically disparate areas. The study defined a Health Disparity Zone (HDZ) as a zipcode-defined area in which diabetes prevalence was above the national average of diabetes prevalence among minorities. Within a HDZ, individuals were further subdivided by zip code areas, depending upon whether the zip code area contained more than 50 percent minorities or not (minority vs. nonminority). The study defined a non-HDZ as one with diabetes prevalence at or below the national average for minorities. The study population thus included 3359

individuals residing in HDZs and 34,066 individuals residing in non-HDZs. Of the 3359 members living in HDZs, 2068 (61.6 percent) lived within minority zip code areas.

The study assessed the degree to which a patient reminder telephone intervention increased HbA1c testing rates among non-adherent members, and analyzed differences by zone and minority neighborhood status.<sup>32</sup> Non-adherent members were defined as those who lacked an HbA1c test in the baseline period.

After 1 year, the rate of testing increased significantly (p<0.0001) in the HDZ group to 59.4 percent. The rate of testing also increased significantly by 4.6 percent to 68.6 percent in the non-HDZ group (p<0.0001). Thus, the initial 12 percent disparity in testing between high disparity and low disparity zones was narrowed by 3 percent to 9 percent after 1 year. The 3 percent reduction in disparity did not reach standard statistical significance by Fisher's exact two-tailed test (p=0.06). \*\*\*

The authors further examined the potential for a differential effect in minority neighborhoods compared to non-minority neighborhoods within the high disparity zone. Rates of testing in zip codes with greater numbers of minority members increased significantly from baseline to 1 year, with an increase in testing of 15.5 percent (p< 0.0001). Conversely, no increase over time was observed in nonminority zip codes in the HDZ. Therefore, a reduction in racial disparity was noted within the population with high prevalence of diabetes, while no reduction in disparity between high and low prevalence zones overall was seen.

Because there was no unexposed group in this study, it is not possible to be certain that observed increases in HbA1c testing in both high disparity and non disparity zones were due to intervention effects. Nonetheless, a racial disparity was decreased within the high disparity zone, where minority zip codes demonstrated greater increases in testing rates compared to non-minority zip codes. Whether this was because it is easiest to "move the dial" in populations that start with the poorest health measures is unknown. Nonetheless, in a "per-protocol" analysis by whether or not the call was actually received, receipt of the phone reminder was associated with increased likelihood of HbA1c testing, suggesting that phone reminders targeted to disadvantaged populations may be a tool for further evaluation.

One study measured lipid testing as the outcome.<sup>31</sup> This prospective cohort study used a database of Medicare patients in New York to evaluate an intensive intervention program designed to reduce the recognized racial disparity in the rate of biennial lipid profile testing among New York City residents with diabetes between 1999 and 2004.<sup>31</sup>

The study was quasi-experimental, of good quality, and reported both a before-and-after analysis for the African-American Medicare fee-for-service (FFS) beneficiaries with diabetes living in the Bronx, Kings, New York, and Queens counties, and a comparison against a reference group of all White Medicare FFS beneficiaries with diabetes in the same counties. All subjects had diabetes and received care at physician offices, outpatient clinics, and community health centers in the New York City metropolitan area. The QI intervention compared Medicare New York State Quality Improvement Organization (IPRO) multifaceted provider and community "intense" interventions versus statewide interventions that IPRO implemented for the Physician Office Quality Improvement Project. The outcome of interest was the proportion of beneficiaries with diabetes who received a biennial lipid profile test.

The before (April 1999) and after (March 2002 up to March 2004) comparison reported a 26.2 percent improvement in biennial lipid profile testing rate among African-American beneficiaries from 63.8 percent to 80.5 percent and an 8.3 percent improvement in biennial lipid profile testing rate among White beneficiaries (increase from 82.8 percent to 89.7 percent).<sup>31</sup>

<sup>\*\*\*</sup>Fisher's Exact Two-Tailed Test, calculated by review authors using reported data.

The baseline disparity in the biennial lipid profile testing rate among African-American beneficiaries compared with all eligible White beneficiaries in the intervention areas was 19 percent. The disparity during the 2002 to 2004 retesting period was 9.2 percent, a reduction of 52 percent. The "intervention" itself was multi-faceted, targeting individuals and providers in many settings and with different specific approaches, including a provider toolkit, patient educational materials, provider reminders, on-site visits and cultural competency training. At the same time, community-directed interventions were put into place including outreach, focus groups and community-based education. As a community-level intervention, no attempt was made to assess individual receipt of the intervention itself; rather, the outcomes were measured at the population level. Thus, individuals could move in and out of the population. Nonetheless, a reduction in disparity was observed over the intervention period, and despite the fact that it cannot be directly associated with the intervention itself or to components of the complex intervention, intermediate goals such as participation in cultural competency training were met.<sup>31</sup>

# **Effects on Insurance Disparities**

### Overview

Two studies examined a difference in outcomes associated with insurance status (Table 5). Both evaluated interventions to improve cancer screening rates. In one study, <sup>19</sup> a patient reminder system was equally successful in commercially insured and Medicare insured women at increasing rates of regular mammography screening. The study was conducted in a large group–model health maintenance organization (HMO) and assessed the use of patient reminders, including a mailed reminder (n=630) or telephone contact with opportunity to schedule screening (n=653) as compared with usual care (n=625) among women aged 50 to 75 years who had received a previous mammogram but were overdue for breast cancer screening. <sup>19</sup>

The second study comprised a secondary analysis of an RCT conducted in a large primary care practice-based research network, aimed at increasing colorectal cancer (CRC) screening among eligible adults and including 153 individuals with private insurance and 312 with public insurance.<sup>26</sup>

Table 5. Summary of effects on disparities in health outcomes associated with insurance status

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Barr et al., 2001 <sup>19</sup> RCT Breast cancer screening	Patient reminder system	<ul> <li>No disparity was observed at baseline.</li> <li>The intervention was successful in both groups.</li> <li>Reminder interventions improved likelihood of screening mammography in both commercially insured women (p=0.001) and women covered by Medicare (p=0.01) with no difference in improvement between groups.</li> </ul>
Lasser et al., 2011 <sup>26</sup> RCT Colorectal cancer screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in screening rates at baseline by race or ethnicity.</li> <li>The intervention increased screening rates in both the private and public insurance groups compared to individuals in the usual care group.</li> <li>The intervention was associated with a better screening rate for the privately insured group compared to publicly insured group.</li> </ul>

**Abbreviations:** QI = quality improvement; RCT = randomized controlled trial.

### **Detailed Description**

A subgroup analysis explored variable effects of the QI strategies in commercially insured patients as compared with Medicare patients, representing approximately 96 percent of the original trial population. The quality of this study was assessed as poor. Among commercially insured women, the use of followup mammography was greater among those randomized to telephone contact (55.6 percent), compared with the mail reminder (44.4 percent) or usual care (39.4 percent, p=0.001). Similar effects were observed among women receiving Medicare, who also had greater use of followup mammography when randomized to telephone contact (56.4 percent), compared with the mail reminder (43.6 percent) or usual care (42.7 percent, p=0.01).

Women covered by Medicare had the same likelihood of followup mammography during the study as commercially insured women. Across insurance types, the difference in followup mammography was not significantly different with the mailed reminder, compared with usual care (p=0.25). Thus, compared with usual care, the telephone based intervention improved mammography use for all populations, but the mailed reminders did not. The telephone based intervention did not demonstrate greater effect in disadvantaged women, as women in the commercially insured and Medicare groups derived similar benefit.

The other subgroup analysis involved an RCT in which patients due for CRC screening were assigned to either usual care or language-concordant assistance by a patient navigator, a trained individual who promoted self-management strategies and provided patient education and reminders. The intervention as compared with usual care was associated with significantly increased screening rates among both privately insured (43.4 percent vs. 22.1 percent, p=0.005) and publicly insured participants (28.9 percent vs. 18.9 percent, p=0.04), though the effectiveness of the navigator intervention appeared somewhat increased in the privately insured group during the 12-month followup period.<sup>26</sup>

## **Effects on Language Disparities**

### Overview

Two studies examined the degree to which language concordance was associated with increased cancer screening (breast and colorectal) among individuals speaking primarily English or other languages (Table 6). Both of them studied language concordance, in which strategies are provided in the native or preferred language of the participant (e.g., in Spanish for native Spanish speakers).

One study was a secondary analysis of an RCT which provided language-concordant patient education for English speaking and Spanish speaking participants, with the goal of increasing adherence to recommendations for breast and colorectal cancer screening among eligible adults. For breast cancer screening, Spanish speakers were more likely to be up to date at baseline than English speakers (OR 1.46; 95% CI: 1.16 – 1.84). The intervention was associated with increased rates of screening overall, with subgroup analysis indicating a greater effect in the Spanish speaking group (OR 1.85; 95% CI: 1.38, 2.47) than the English speaking group (OR 1.18; 95% CI: 0.82, 1.71). However, the overall multivariate analysis failed to confirm these results and providing the intervention in Spanish to Spanish speakers did not make it any more effective in this group. This may suggest that targeted language concordant interventions could warrant further examination, but these results reflect only one study. For colorectal screening, there was no difference in up to date status at baseline, the intervention was again effective overall, and there was no language-by-intervention effect.

Another secondary analysis of an RCT included language-concordant assistance by a patient navigator who promoted self-management strategies and provided patient education and reminders to facilitate adherence to colorectal cancer screening among eligible adults in a large primary care practice-based research network, including 224 individuals speaking English as their primary language and 241 speaking a language other than English, including Spanish, Portuguese, and Haitian Creole. <sup>26</sup>

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Beach et al., 2007 <sup>20</sup> RCT Cancer: CRC and breast cancer screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>At baseline, Spanish speakers were more likely to be up to date on breast cancer screening.</li> <li>The intervention was effective at increasing rates of breast cancer screening overall, with greater effect among Spanish speakers.</li> <li>Nonetheless, the difference between observed effects for breast cancer screening in the two language groups was not significant.</li> <li>No disparity in CRC screening rate was measured at baseline.</li> <li>The intervention was associated with increases in CRC screening in both groups, with neither group associated with a greater effect of the intervention.</li> <li>Therefore, although there was no evidence that the intervention might reduce known disparities, the intervention was effective at increasing CRC screening for both groups.</li> </ul>
Lasser et al., 2011 <sup>26</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self- management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity was measured at baseline</li> <li>English speaking participants had a similar incidence of CRC screening during 1 year of followup in the intervention group as compared with usual care.</li> <li>Intervention was particularly beneficial for non-English language participants</li> </ul>

**Abbreviations:** CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial.

# **Detailed Description**

In one fair RCT<sup>20</sup>, a prevention care manager periodically called to remind women about screening, assist in overcoming barriers, provide emotional support, and help with scheduling; women also received additional educational materials in their preferred language by mail. A subgroup analysis focused on the impact of this strategy among women who preferred Spanish (n=848) and those who preferred English (n=498). Among Spanish-speaking women, a significantly greater proportion of women (72 percent) in the care management group were up to date on breast cancer screening at followup, compared with 58 percent in those receiving usual care, (adjusted OR 1.86, 95% CI, 1.39 to 2.50). Among English speakers, women in the care management group were not significantly more likely to be up to date on breast screening, compared with women in the usual care group (60 percent and 56 percent, respectively; adjusted OR 1.23, 95% CI, 0.85 to 1.78). Although Spanish-speaking women benefited more from care management than did English-speakers, when assessed statistically as an interaction effect, the result was non-significant (OR 1.51, 95% CI, 0.94 to 2.42). Though women speaking either language were equally likely to receive appointment reminders, access advice, and assistance with making primary care appointments, Spanish-speaking women were more likely than English–speaking women to receive help scheduling appointments for breast cancer screening (26.5 vs. 18.4 percent, p=0.01) and were more likely to have educational materials mailed to them (70.2 vs. 60.6 percent, p=0.01). Although it was not significantly greater than the effect seen in English-speakers, the magnified effect observed in Spanish-speakers may suggest that language concordance is helpful in interventions seeking to achieve up to date cancer screening status.

Another secondary RCT analysis examined whether education and reminders from a language concordant patient navigator led to differences in CRC screening participation among English speakers as compared with non-English speakers. <sup>26</sup> While English speaking participants had a similar incidence of CRC screening during 1 year of followup in the intervention group as compared with usual care (26.8 percent vs. 21.4 percent, p=0.35), assistance from a patient navigator was associated with increased CRC screening among individuals speaking languages other than English as their primary language (28.9 percent vs. 18.9 percent, p=0.04). <sup>26</sup>

# **Effects on Health Literacy Disparities**

#### Overview

In one study focusing on improving provider-patient communication, in the VA system, colorectal cancer screening was increased among individuals with limited health literacy (55.7 percent vs. 30 percent; p=.002) but not among individuals with adequate health literacy (39 percent vs. 36 percent; p=.65) in the 20 percent subsample that underwent literacy assessment. The intervention itself included a workshop and feedback sessions for providers and educational materials for patients that included a video (Table 7).

Table 7. Summary of effects on disparities in health outcomes associated with health literacy

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Ferreira et al., 2005 <sup>23</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Provider education</li> <li>Audit and feedback</li> </ul>	<ul> <li>Patients with limited health literacy were significantly more likely to be screened for CRC when treated at the VA clinic implementing the QI strategy as compared with patients treated at the usual care clinic (55.7% vs. 30.0%, p=0.002).</li> <li>Patients with adequate health literacy were equally likely to pursue CRC screening when treated at the VA clinic implementing the QI strategy as compared with the patients treated at the usual care clinic (39.0% vs. 36.0%, p=0.65).</li> <li>Although the effect of the intervention on disparity was not measured directly, the intervention improved the incidence of up-to-date CRC screening among those with limited health literacy only and not among those with higher health literacy, suggesting that it might be a useful tool for reducing literacy related disparity.</li> </ul>

**Abbreviations:** CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial; VA = Veterans Affairs.

# **Detailed Description**

The RCT included men aged 50 years or older, without a family history of CRC or polyps, who had not received recent CRC screening. The study randomized one VA clinic to a provider and patient education strategy and one VA clinic to continue usual care. <sup>23</sup> The provider education intervention included a two-hour workshop on CRC screening and health literacy—sensitive communication skills, 1-hour feedback sessions every 4 to 6 months on the clinic's CRC screening rates, confidential provider—specific reports on screening rates, and small group role—playing about how to effectively make CRC screening recommendations. The patient education intervention included a video and educational material on CRC screening, though only

204 of 1978 patients in the overall RCT received this. The quality of this study was assessed as poor; randomization was broken, and the participants in this analysis represent a small proportion of the entire data set. A subset of patients completed the Rapid Estimate of Adult Literacy in Medicine (REALM) instrument at baseline, representing approximately 19 percent of the overall study sample (n=382/1978). Patients were classified as limited health literacy if they scored below ninth grade level on the REALM (n=139) and adequate health literacy if they scored at the ninth grade level or above (n=243). Patients with limited health literacy who were treated at the intervention clinic were significantly more likely to undergo CRC screening (55.7 percent) than low literacy patients treated at the usual care clinic (30.0 percent, p=0.002). There was no significant difference in CRC screening between the intervention and usual care clinics for patients with high health literacy (39.0 vs. 36.0 percent, p=0.65). Although the effect of the intervention on disparity was not measured directly, the intervention improved the incidence of up-to-date CRC screening among those with limited health literacy only and not among those with higher health literacy, suggesting that it might be a useful tool for reducing literacy related disparity.

## **Effects on Socioeconomic Disparities**

#### Overview

In the two studies that assessed differences in effect by socioeconomic status, no effect was seen by income, but individuals with less education experienced greater benefits of collaborative care for depression than did those with higher education (Table 8).

Table 8. Summary of effects on disparities in health outcomes associated with socioeconomic status

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Arean et al., 2007 <sup>18</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (collaborative care model)</li> <li>Facilitated relay of clinical data to provider</li> </ul>	Both low income and high/middle income populations experienced a very small benefit from the collaborative care intervention [fewer depression symptoms (high/middle income adjusted OR -0.41, 95% CI -0.49 to -0.33; low-income adjusted OR -0.39, 95% CI, -0.5 to -0.27; comparator: usual care)], but no disparities in depressive symptoms had existed at baseline.
Bao et al., 2011 <sup>24</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (collaborative care model)</li> <li>Provider reminder system</li> </ul>	<ul> <li>No disparity in depressive symptoms was present at baseline</li> <li>At 24 months, participants with no college education had a greater reduction in depression than participants with college.</li> </ul>

**Abbreviations:** OR = odds ratio; QI = quality improvement; RCT = randomized controlled trial.

## **Detailed Description**

The IMPACT trial on depression care <sup>17,18</sup> which also examined racial or ethnic disparities (see description above), included a subgroup analyses to evaluate whether this intervention could decrease outcome disparities among older ethnic minorities <sup>17</sup> and among older adults of higher and lower incomes. <sup>18</sup> Both low income and high/middle income populations experienced a very small benefit from the collaborative care intervention [fewer depression symptoms (high/middle income adjusted OR -0.41, 95% CI -0.49 to -0.33; low income adjusted OR -0.39, 95% CI, -0.5 to -0.27; comparator: usual care)], but no disparities in depressive symptoms had existed at baseline. <sup>18</sup>

A functional impairment gap did exist at baseline, with poor individuals scoring lower on the PCS ( $41.1 \pm 7.4$  vs.  $38.6 \pm 7.1$ ; p<0.01). At 1 year, the collaborative care groups did experience a small but nonsignificant benefit when compared with usual care groups across income levels (high/middle income: adjusted OR 1.67, 95% CI, 0.78 to 2.55; low income: adjusted OR 1.46, 95% CI, 0.33 to 2.60), the effect of which was to reduce the gap somewhat between groups. Nonetheless the intervention was not associated with improvements overall. <sup>18</sup>

The PROSPECT study on depression care, <sup>24</sup> which also examined race or ethnicity disparities (see description above) included a subanalysis focused on socioeconomic outcomes, based on degree of college education as a marker for SES. The no-college and some college groups had no statistically significant difference in depression severity as measured by the Hamilton Depression Rating Scale, at baseline or up to 18 months post-intervention, although at 24 months, the no-college group reduced their depression scores more than did the group with a college education (difference of -3.8 [-6.8 to -0.4]). <sup>24</sup> Nonetheless, no disparity existed to be narrowed, and the authors do not provide data to attribute the effect to the intervention. After adjusting for baseline antidepressant use, the no-college educated recipients of the study intervention had greater benefit in appropriate use of antidepressants (i.e., adequate dose) compared with the college-educated participants who received intervention (23.4 percent difference, 95% CI, 5.5 to 43.7 percent) at 1 year, but a statistically significant difference was not sustained at 24 months.

# **Effects on Disparities by Sex**

### Overview

One analysis from the PIC depression study examined the degree to which a collaborative care model for depression could reduce known disparities in accessing care and in outcomes by sex (Table 9). The intervention components: (1) enrolled practices provided in-kind resources (up to one half of the cost) to assist in intervention implementation; (2) provider training at study onset and monthly meeting thereafter to provide feedback on treatment patterns; (3) each practice had a designated staff nurse who was trained as a "depression specialist" who would screen patients for depression, then educate and engage those who screened positive in depression self-management. The QI–Meds intervention contained all of the above components plus trained nurses who assessed patient adherence to antidepressant pharmacotherapy. The QI–Therapy intervention contained all of the above components plus local psychotherapists who could provide individual or group cognitive behavioral therapy.

Women had higher rates of appropriate depression care compared with men at 2 years (p = 0.0001). A medication focused intervention and a therapy focused intervention decreased a

disparity gap between men and women in probable unmet need from 10 percent to 1 percent (QI–Meds) and 3 percent (QI–Therapy) at 24 months.

Table 9. Summary of effects on disparities in health outcomes associated with sex

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Sherbourne et al., 2004 <sup>30</sup> Cluster RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>Women had higher rates of appropriate depression care compared with men at 2 years (p= 0.0001).</li> <li>QI-meds and QI-Therapy decreased "probable unmet need" disparity gap between men and women from 10% to 1% (QI-Meds) and to 3% (QI-Therapy) at 24 months.</li> </ul>

**Abbreviations:** QI = quality improvement; RCT = randomized controlled trial.

## **Detailed Description**

Women in QI–Meds group had non-sustained increase in employment rates compared with usual care women (p=0.23); no change occurred among women in QI–Therapy compared with usual care (p=0.10). Men in QI–Therapy also had a non-sustained improvement in employment status compared with the usual care (p=0.61) and QI–Meds (p=0.71) groups. The employment status disparity between men and women widened at 2 years in both intervention groups (QI–Meds gap = 8 percent; QI–Therapy gap = 6 percent) with men less likely to be employed than women.  $^{30}$ 

Women were 10.2 percent more likely than men at baseline to receive appropriate depression care (p=0.03). Over time, the disparity gap for "probable unmet need for depression care" between men and women was decreased from 10 percent to 1 percent (QI–Meds) and to 3 percent (QI–Therapy) at 2 years.<sup>30</sup>

### **Discussion**

## **Review of Main Findings**

Disparities in health care access, utilization and outcomes are a known challenge to improving quality of care across the United States. Disparities have been measured in a wide range of clinical conditions including: cancer, diabetes, end stage renal disease, cardiovascular disease, HIV and AIDS, maternal and child health, mental health and substance abuse, and respiratory diseases. Given the potential for quality improvement (QI) strategies to improve health care in clinical settings, interest has developed in whether they might be used to reduce disparities, potentially by being particularly effective at improving outcomes in individuals and groups affected by disparities. A review of effectiveness of QI interventions to reduce disparities has the potential to provide a basis for policy and health care program decision making when choices about how to best serve a disadvantaged population are made.

QI activities that are implemented across a large population provide a degree of efficiency and sustainability. However, even in the face of positive outcomes, disparities remain, as improvements in care affect all segments of the population—the "rising tide raises all boats." Conversely, it is theoretically possible that QI interventions could worsen quality of care for a disadvantaged population relative to an advantaged one.

Although substantial research has examined ways to improve the health of specific populations known to have disproportionate rates of poor outcomes, little research has focused on measuring a decrease in disparity between two groups. Rather, disparities research generally measures an increase in appropriate care in a disadvantaged group under the assumption that an improvement in outcomes in that group should bring their outcomes closer in line to those of the group to whom they are implicitly compared. This is not always the case. For example, although survival has improved for premature infants overall in the United States, including among racial minorities, the gap between survival for White and Black babies has continued to increase. Thus, research may focus on changing outcomes in one group, but if improvements are seen in all groups, disparities will remain. The purpose of this review was specifically to identify the potential for QI interventions to close the gap between measured outcomes in two groups of individuals to identify interventions that might be specifically targeted to situations where there is a disparity.

Methodologic challenges in this literature, including heterogeneity in the study populations, target clinical conditions, and interventions have made synthesis and interpretation challenging. No two studies used the same QI intervention to study disparity in the same health outcome. The standard approach to assessing the strength of an overall body of literature involves assessing a number of factors, including the risk of bias in individual studies, consistency of the overall literature, and precision of available estimates. The resulting strength of evidence score reflects the degree to which we expect the observed effects to be stable and not altered with future research. For this particular review, the measure of effect was a change in a disparity between two groups in a health outcome, rather than a change in the health outcome itself. The strength of the evidence would nonetheless be assessed for each health outcome and each intervention. Thus, as the results could not be combined quantitatively, and no two studies evaluated the same intervention and outcomes, it is not possible to consider the strength of the evidence (confidence in a given effect) to be other than insufficient at this time.

Nonetheless, 19 papers qualified for our review, reflecting results from 14 studies, and these studies do provide information on the potential role of some QI interventions in reducing disparities in health and health care, pending additional research. Studies were conducted to improve care in cancer screening, cardiovascular disease, depression, and diabetes care, using a number of different QI models (Table 10). Most interventions included some elements of patient education (12/14) and promotion of self-management (10/14). Most (11/12) involved at least two QI components. One included language concordance as an additional component.<sup>20</sup>

Table 10. Included studies' quality improvement strategies

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Author, Year Clinical Condition(s)	Patient Education	Provider Education	Promotion of Self- Management	Foodback	Facilitated Relay of Clinical Data to Providers	Patient Reminder System	Provider Reminder System	Organizational Change	Other
Arean et al., 2007; <sup>18</sup> Arean et al., 2005 <sup>17</sup> Depression	•	•	•		•				● <sup>a</sup>
Bao et al., 2011 <sup>24</sup> Depression	•	•	•				•		● <sup>a</sup>
Barr et al., 2001 <sup>19</sup> Cancer: breast cancer screening						•			
Beach et al., 2007 <sup>20</sup> Cancer: CRC and breast cancer screening	•		٠			•			
Bosworth et al., 2011 <sup>21</sup> Cardiovascular disease: hypertension	•		•						
Coberley et al., 2007 <sup>32</sup> Diabetes	•		•					• <sup>b</sup>	
Connett and Stamler, 1984 <sup>22</sup> Cardiovascular disease: coronary artery disease and hypertension	•		•						
Ferreira et al., 2005 <sup>23</sup> Cancer: CRC screening	•	•		•					
Lasser et al., 2011 <sup>26</sup> Cancer: CRC screening	•		•			•			

Table 10. Included studies' quality improvement strategies (continued)

Author, Year Clinical Condition(s)	Patient Education	Provider Education	Promotion of Self- Management	Audit and Feedback	Facilitated Relay of Clinical Data to Providers	Patient	Provider Reminder System	Organizational Change	Other
Mahotiere et al., 2006 <sup>31</sup> Diabetes	•	•	•	•					•°
Olomu et al., 2010 <sup>33</sup> Cardiovascular disease: coronary artery disease	•	•	•						●d
Miranda et al., 2003; <sup>15</sup> Miranda et al., 2004; <sup>34</sup> Wells et al., 2007; <sup>28</sup> Wells et al., 2004; <sup>29</sup> Sherbourne et al., 2004 <sup>30</sup> Depression	•	•	•	•	•				● <sup>a</sup>
Sequist et al., 2010 <sup>16</sup> Diabetes		•		•					
Siddiqui et al., 2011 <sup>25</sup> Cancer: CRC screening	•	•				•			
Counts	12	8	10	4	2	4	1	1	5

**Notes:** <sup>a</sup>Collaborative care; <sup>b</sup>Disease Management; <sup>c</sup>Community intervention; <sup>d</sup>Guideline adherence **Abbreviation:** CRC = colorectal cancer.

Although the strength of the evidence (our confidence that future studies will not change observe estimates of effect) is uniformly insufficient at this point in time, this is partly due to the substantial heterogeneity in research clinical targets, disparity targets and interventions. Additional research is needed to confirm currently observed effects in individual studies, and provide new information on approaches in the health system to reducing disparities in health and health care. Therefore, we present the evidence in two ways, by clinical target and by disparity target, to describe areas where there may be promising suggestions for additional research.

## **Effects on Disparities**

Included studies addressed disparities by race or ethnicity (Table 11), insurance (Table 12), language (Table 13), health literacy (Table 14), socioeconomic status (Table 15), and sex (Table 16).

Table 11 summarizes results of studies addressing disparities in health outcomes associated with race or ethnicity. One disease management and patient education program<sup>32</sup> was associated with a reduction in disparity between Black and White patients in HbA1c testing when it was targeted in a health disparity zone (HDZ) with a high prevalence of diabetes. Other interventions did not demonstrate a significant reduction in disparity, but an amplified effect was seen in

nonwhite (presumably disadvantaged) population, including an additional patient education program for reduction in blood pressures, <sup>21</sup> and a complex collaborative care model aimed at providers of patients with depression. <sup>15,28,29,34</sup> In the latter study, the intervention was more effective in the short term among minorities, although the interaction was no longer significant after 1 year, and the intervention was not effective overall at five and nine years.

Table 11. Summary of effects on disparities in health outcomes associated with race or ethnicity

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity
Arean et al., 2005 <sup>17</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (collaborative care model)</li> <li>Facilitated relay of clinical data to provider</li> </ul>	<ul> <li>No disparity in depression severity existed by race or income at baseline.</li> <li>The intervention was effective in all racial subgroups with no interaction by race and no amplified effect in any group.</li> <li>In subgroup analysis, the intervention was associated with greater use of psychotherapy but not pharmacotherapy within the Black population</li> </ul>
Bao et al., 2011 <sup>24</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (collaborative care model)</li> <li>Provider reminder system</li> </ul>	<ul> <li>At baseline within the usual care group, 22% of minorities had adequate antidepressant use, compared to 39% of White patients.</li> <li>The intervention had no effect on this disparity and ethnic minorities did not receive greater benefit from intervention compared with White patients during any time period.</li> </ul>
Bosworth et al., 2011 <sup>21</sup> RCT Cardiovascular disease: hypertension	<ul> <li>Patient education</li> <li>Promotion of self- management</li> </ul>	<ul> <li>The race by time by treatment group effect model suggested differential intervention effects on BP over time for White participants vs. nonwhite participants for both SBP (p=0.08) and DBP (p=0.01).</li> <li>The combination of home BP monitoring and tailored behavioral intervention was most effective in nonwhite participants at 24 months (p=0.04).</li> </ul>
Coberley et al., 2007 <sup>32</sup> Retrospective cohort Diabetes	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Organizational change (disease management)</li> </ul>	<ul> <li>Initial racial disparity in HbA1c testing between the health disparity zone (HDZ) group and non-HDZ group was 12%.</li> <li>Disparity was not significantly reduced after 12 months (p=0.06).</li> <li>Within the HDZ zone (high prevalence of diabetes), testing increased by15% among Black participants but not among White participants, resulting in a reduction in disparity in this subgroup analysis.</li> </ul>

Table 11. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity			
Connett and Stamler, 1984 <sup>22</sup> RCT Cardiovascular disease: coronary artery disease and hypertension	<ul> <li>Patient education</li> <li>Promotion of self-management</li> </ul>	<ul> <li>At baseline, Black participants had higher rates of smoking than White participants (68.7% vs. 63%; p&lt;.001).</li> <li>Both racial groups experienced significant reductions in smoking of close to 50% in the intervention group and more than 35% in the usual care group.</li> <li>The baseline disparity persisted in the intervention group but was apparently reduced in the usual care group.</li> <li>A statistically significant but clinically insignificant disparity in DBP and SBP by race was present at baseline.</li> <li>Blood pressures were reduced in both the intervention and control groups, with greater change observed in the intervention group.</li> <li>The small disparity observed at baseline was further reduced at followup in the intervention group but not the control group.</li> </ul>			
Lasser et al., 2011 <sup>26</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates at baseline by race or ethnicity.</li> <li>The intervention was more effective in White and Black individuals relative to those of other or unknown race.</li> </ul>			
Mahotiere et al., 2006 <sup>31</sup> Prospective cohort Diabetes	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Other (community intervention)</li> </ul>	<ul> <li>The disparity in biennial lipid profile testing at baseline was 19%.</li> <li>The biennial lipid profile testing rate improved by 26.2 percent in African-American beneficiaries with diabetes in the intervention areas following implementation of the QI program.</li> <li>The disparity in performance of biennial lipid profile between African-American Medicare fee-for-service beneficiaries and White Medicare fee-for-service beneficiaries reduced to 9.2% following implementation of the QI program.</li> <li>An analysis of the direct impact of the selected interventions on reducing the disparity in this uncontrolled database analysis was not feasible.</li> </ul>			

Table 11. Summary of effects on disparities in health outcomes associated with race or ethnicity (continued)

Author, Year Study Design Clinical Condition(s)	QI Intervention Characteristic(s)	Effect on Health Disparity			
Miranda et al., 2003 <sup>15</sup> ; Miranda et al., 2004 <sup>34</sup> ; Wells et al., 2007 <sup>28</sup> ; Wells et al., 2004 <sup>29</sup> Cluster RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Other (collaborative care model)</li> <li>Facilitated relay of clinical data to provider</li> </ul>	<ul> <li>The intervention was associated with decreases in probable depressive disorder among minorities but not White patients at 12 months. (Latino p=0.02; African-American p=0.01).</li> <li>At 12 months among intervention recipients, the baseline disparity increased (6.7% to 7.7%) between Latino and White patients, and decreased between African-American and White patients (9.2% to 6.7%).</li> <li>Although a statistically significant interaction was seen between intervention and ethnicity at 6 months when minorities were grouped and contrasted with White patients, no such interaction persisted at 12 months.</li> <li>The overall effect of the intervention on depression status was not significant at five and nine years, but an interaction with race was seen in the overall model of effectiveness. The intervention was associated with improvements in the Mental Health Inventory among minorities (p=0.008) but not among White patients (p=0.59).</li> <li>In subanalysis at five years, Ql-Therapy but not Ql-Meds was effective within the minority population.</li> </ul>			
Olomu et al., 2010 <sup>33</sup> Retrospective cohort (historic controls) Cardiovascular disease: coronary artery disease	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Other (guideline adherence)</li> </ul>	<ul> <li>The American College of Cardiology's Acute Myocardial Infarction (AMI) Guidelines Applied in Practice (GAP) strategy was associated with increased inpatient use of beta-blockers among nonwhite patients.</li> <li>Racial disparities in use of cardiac catheterization and percutaneous coronary intervention appeared after implementation of the GAP QI strategy despite overall improvements in care.</li> <li>The admission tool and inpatient aspirin were more often used post-GAP vs. pre-GAP in both White and nonwhite patients.</li> </ul>			
Sequist et al., 2010 <sup>16</sup> Cluster RCT Diabetes	<ul><li>Provider education</li><li>Audit and feedback</li></ul>	<ul> <li>Disparities were present at baseline in HbA1c levels, BP control, and LDL level.</li> <li>The intervention showed no effect overall in either racial group.</li> <li>The intervention did not reduce the disparity.</li> </ul>			
Siddiqui et al., 2011 <sup>25</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates at baseline by race or ethnicity.</li> <li>No statistically significant difference in screening rates between Whites and African-Americans in the control group.</li> <li>When intervention groups were combined, the screening rate was significantly higher in Whites compared to African-Americans.</li> </ul>			

**Abbreviations:** BP = blood pressure; CRC = colorectal cancer; DBP = diastolic blood pressure; GAP = American College of Cardiology's AMI Guidelines Applied in Practice; HbA1c = hemoglobin A1c; HDZ = health disparity zone; LDL = low density lipoprotein; QI = quality improvement; RCT = randomized controlled trial; SBP = systolic blood pressure.

Two studies (Table 12) examined a difference in outcomes associated with insurance status. <sup>19,26</sup> In both studies, the intervention was equally successful at increasing cancer screening

in publicly and privately insured participants. In the first, <sup>19</sup> a patient reminder system for breast cancer screening improved mammography rates in all women. In the second study, <sup>26</sup> language-concordant assistance by a patient navigator who promoted self-management strategies, patient education, and reminders was associated with significantly increased colorectal cancer screening rates among both privately insured and publicly insured participants compared to usual care, but seemed to be most effective in the privately insured group.

Table 12. Summary of effects on disparities in health outcomes associated with insurance status

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
Barr et al., 2001 <sup>19</sup> RCT Cancer: breast cancer screening	Patient reminder system	<ul> <li>No disparity was observed at baseline.</li> <li>The intervention was successful in both groups.</li> <li>Reminder interventions improved likelihood of screening mammography in both commercially insured women (p=0.001) and women covered by Medicare (p=0.01) with no difference in improvement between groups.</li> </ul>
Lasser et al., 2011 <sup>26</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity in CRC screening rates at baseline by race or ethnicity.</li> <li>The intervention increased screening rates in both the private and public insurance groups compared to individuals in the usual care group.</li> <li>The intervention was associated with a better screening rate for the privately insured group compared to publicly insured group.</li> </ul>

**Abbreviations:** CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial.

Two studies examined the effects of QI strategies on disparities associated with language (Table 13). Both of them studied language concordance, in which strategies are provided in the native or preferred language of the participant (e.g., in Spanish for native Spanish speakers). One study examined the degree to which a language concordant intervention increased cancer screening (breast and colorectal) among English and Spanish speaking patients. For breast cancer screening, Spanish speakers were more likely to be up to date at baseline than English speakers (OR 1.46; 95% CI: 1.16 – 1.84). The intervention was associated with increased rates of screening overall, with greater effect in the Spanish speaking group (OR 1.85; 95% CI: 1.38, 2.47) than the English speaking group (OR 1.18; 95% CI: 0.82, 1.71). However, the overall multivariate analysis failed to confirm these results and providing the intervention in Spanish to Spanish speakers did not make it any more effective in this group. For colorectal screening, there was no difference in up to date status at baseline, the intervention was again effective overall, and there was no language-by-intervention effect.

A second study included language-concordant assistance by a patient navigator promoting self-management strategies, and providing patient education and reminders to facilitate adherence to colorectal cancer screening for individuals speaking English as their primary language and individuals speaking a language other than English. The patient navigator intervention was associated with increased CRC screening among individuals speaking languages other than English as their primary language (28.9 percent vs. 18.9 percent, p=0.04), but not among patients for whom English was their primary language (26.8 percent vs. 21.4 percent, p=0.35)<sup>26</sup> These studies combined may suggest that targeted language concordant interventions could warrant further examination, with results suggesting a significantly different

effect for non-English speakers and English speakers in one study, and a clinically, but not statistically different effect in the other.

Table 13. Summary of effects on disparities in health outcomes associated with language barrier

Author, Year Study Design Clinical Condition	dy Design Characteristic(s) Effect on Health Dispa			
Beach et al., 2007 <sup>20</sup> RCT Cancer: CRC and breast cancer screening	<ul> <li>Patient education</li> <li>Promotion of self-management</li> <li>Patient reminder system</li> </ul>	<ul> <li>At baseline, Spanish speakers were more likely to be up to date on breast cancer screening.</li> <li>The intervention was effective at increasing rates of breast cancer screening overall, with greater effect among Spanish speakers.</li> <li>Nonetheless, the difference between observed effects for breast cancer screening in the two language groups was not significant.</li> <li>No disparity in CRC screening rate was measured at baseline.</li> <li>The intervention was associated with increases in CRC screening in both groups, with neither group associated with a greater effect of the intervention.</li> <li>Therefore, although there was no evidence that the intervention might reduce known disparities, the intervention was effective at increasing CRC screening for both groups.</li> </ul>		
Lasser et al., 2011 <sup>26</sup> RCT Cancer: CRC screening	<ul> <li>Patient education</li> <li>Promotion of self- management</li> <li>Patient reminder system</li> </ul>	<ul> <li>No disparity was measured at baseline</li> <li>English speaking participants had a similar incidence of CRC screening during 1 year of followup in the intervention group as compared with usual care.</li> <li>Intervention was particularly beneficial for non-English language participants</li> </ul>		

**Abbreviations:** CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial.

In one health literacy study focusing on improving provider-patient communication in the VA system, colorectal cancer screening was increased among individuals with limited health literacy (55.7 percent vs. 30 percent; p=.002) but not among individuals with adequate health literacy (39 percent vs. 36 percent; p=.65) in the 20 percent subsample that underwent literacy assessment. The intervention itself included a workshop and feedback sessions for providers and educational materials for patients that included a video (Table 14).

Table 14. Summary of effects on disparities in health outcomes associated with health literacy

<ul> <li>Patient education         <ul> <li>Postient education</li> <li>Provider education</li> <li>Audit and feedback</li> </ul> </li> <li>Patients with limited health literacy were significantly more likely to be screened for CRC when treated at the VA clinic implementing the QI strategy as compared with patients treated at the usual care clinic (55.7% vs. 30.0%, p=0.002).</li> <li>Patients with adequate health literacy were equally likely to pursue CRC screening when treated at the VA clinic implementing the QI strategy as compared with the patients treated at the usual care clinic (39.0% vs. 36.0%, p=0.65).</li> <li>Although the effect of the intervention on disparity was not measured directly, the intervention improved the incidence of up-to-date CRC screening among those with limited health literacy, suggesting that it might be a useful tool for reducing literacy related disparity.</li> </ul>	Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Health Disparity
	RCT	<ul> <li>Provider education</li> </ul>	more likely to be screened for CRC when treated at the VA clinic implementing the QI strategy as compared with patients treated at the usual care clinic (55.7% vs. 30.0%, p=0.002).  • Patients with adequate health literacy were equally likely to pursue CRC screening when treated at the VA clinic implementing the QI strategy as compared with the patients treated at the usual care clinic (39.0% vs. 36.0%, p=0.65).  • Although the effect of the intervention on disparity was not measured directly, the intervention improved the incidence of up-to-date CRC screening among those with limited health literacy only and not among those with higher health literacy, suggesting that it might be a

Abbreviations: CRC = colorectal cancer; QI = quality improvement; RCT = randomized controlled trial; VA = Veterans Affairs.

In the two studies that assessed differences in effect by socioeconomic status, no effect was seen by income, but individuals with less education experienced greater benefits of collaborative care for depression than did those with higher education (Table 15).

Table 15. Summary of effects on disparities in health outcomes associated with socioeconomic status

Status		
Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)	Effect on Disparity Target
Arean et al., 2007 <sup>18</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	Both low income and high/middle income populations experienced a very small benefit from the collaborative care intervention [fewer depression symptoms (high/middle income adjusted OR -0.41, 95% CI -0.49 to -0.33; low income adjusted OR -0.39, 95% CI, -0.5 to -0.27; comparator: usual care)], but no disparities in depressive symptoms had existed at baseline.
Bao et al., 2011 <sup>24</sup> RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Provider reminder system</li> <li>Other (collaborative care model)</li> </ul>	<ul> <li>No disparity in depressive symptoms was present at baseline.</li> <li>At 24 months, participants with no college education had a greater reduction in depression than participants with college.</li> </ul>

**Abbreviations:** OR = odds ratio; QI = quality improvement; RCT = randomized controlled trial.

One analysis examined the degree to which a collaborative care model for depression could reduce known disparities in accessing care and in outcomes by sex (Table 16). Women had higher rates of appropriate depression care compared with men at 2 years (p= 0.0001). A medication focused intervention and a therapy focused intervention decreased a disparity gap

between men and women in probable unmet need from 10 percent to 1 percent (QI–Meds) and 3 percent (QI–Therapy) at 24 months.

Table 16. Summary of effects on disparities in health outcomes associated with sex

Author, Year Study Design Clinical Condition	QI Intervention Characteristic(s)		Effect on Health Disparity
Sherbourne et al., 2004 <sup>30</sup> Cluster RCT Depression	<ul> <li>Patient education</li> <li>Provider education</li> <li>Promotion of self-management</li> <li>Audit and feedback</li> <li>Facilitated relay of clinical data to provider</li> <li>Other (collaborative care model)</li> </ul>	•	Women had higher rates of appropriate depression care compared with men at 2 years (p= 0.0001). QI–Meds and QI–Therapy decreased "probable unmet need" disparity gap between men and women (from 10% to 1% (QI–Meds) and to 3% (QI–Therapy) at 24 months).

**Abbreviations:** QI = quality improvement; RCT = randomized controlled trial.

In sum, there are individual studies that suggest benefits in particular subgroups known to suffer from disparities in health and health care. By far, the largest proportion of the literature focused on the ability of QI interventions to reduce racial disparities, with some suggestions that targeted programs could have some greater effects among racial minorities in both diabetes<sup>32</sup> and hypertension. However, consistent findings from a rich body of literature are lacking to guide QI efforts specifically to reduce disparities.

Based on the limited evidence qualified for inclusion in this report, several strategies are worthy of future study, and possibly wider implementation. These strategies include the collaborative care model 15,17,18,28,29,34 and the role of targeted patient education, 21,26,32 which is accommodating of recipients' language and literacy levels. The collaborative care model emphasizes the coordination of care by multiple clinical providers, usually by a care manager, and interventions that reflect literacy and language needs of participants have included both the preparation of materials in multiple languages, and materials that reflect the complexity of medical or health care information and the need to make it accessible to a range of patients. A common thread is the emphasis on adapting interventions to the needs of disadvantaged populations and to individuals within them. Sufficient data are not available to support universal implementation of these strategies, but the strategies may be suitable for implementation if an appropriate plan is in place to monitor their effectiveness.

# **Applicability**

Given the insufficient strength of evidence health systems or clinicians wishing to replicate any of these interventions should carefully assess whether the interventions apply or must be modified to suit their particular patient population, clinical setting, and available resources. Discussions of implementation are likely to be in the context of specific clinical conditions, given the degree to which the health system is structured in this way. Therefore, we summarize the available literature and its likely applicability by clinical focus below.

#### Cancer

## **Summary**

Five RCTs with subgroup analyses explored the effects of various QI strategies on health care disparities in cancer screening, including one examining breast cancer screening, <sup>19</sup> three studies assessing screening for colorectal cancer, <sup>23,25,26</sup> and one assessing both breast and colorectal cancer screening. <sup>20</sup>. Disparities that served as the focus of these analyses included insurance status, <sup>19</sup> health literacy, <sup>23</sup> and language. <sup>20</sup> The QI strategies included provision of mail or telephone reminders to patients, <sup>19</sup> education and feedback for clinicians, <sup>23</sup> and language—concordant telephone support calls from prevention care managers to patients. <sup>20</sup> All three RCTs took place in study settings including one large group—model health maintenance organization (HMO), <sup>19</sup> two Department of Veterans Affairs (VA) clinics, <sup>23</sup> and 11 community health centers. <sup>20</sup> All studies employed an internal, "usual care" comparison group. A language—concordant intervention, <sup>20</sup> as compared with usual care, was more effective in increasing breast cancer screening among Spanish-speaking women than in English-speaking women, but the observed difference between the two groups (English and Spanish speaking) was not significant. The language-concordant intervention did not have a similar effect on colorectal cancer (CRC) screening. <sup>20</sup> A health-literacy targeted strategy, as compared with usual care, facilitated CRC screening among those with limited health literacy more effectively than among those with high health literacy. <sup>23</sup> A reminder intervention for breast cancer screening had no differential effect on mammography disparities by insurance status. <sup>19</sup>

# **Applicability**

Studies included patients cared for at community clinics in New York City, men treated at two VA clinics in Chicago, and women enrolled in a large group model HMO in the northeastern United States. These settings were appropriate for cancer screening interventions, as the bulk of cancer screening recommendations focus on the clinic setting. However, it is uncertain how well the results of these studies generalize to other populations or settings.

The tested interventions varied substantially, ranging from patient reminders to provider education with audit and feedback. These interventions could be replicated, though they generally required significant organizational resources to develop and implement and may not be feasible in other settings. Barriers to care may also differ in other settings, and the interventions likely would need to be adapted to the needs of the target population. In each study, usual care served as the comparator, and this too, may differ in other practice settings. Thus, the marginal benefit of each intervention likely would be different in different settings.

Study outcomes consisted only of short-term process measures—receipt of cancer screening during followup. No long-term outcomes or clinical outcomes, such as diagnosis of malignancies, were reported. Thus the long-term clinical impact of such interventions is unclear.

## Cardiovascular Disease

# **Summary**

One post hoc analysis of an RCT<sup>22</sup> and one retrospective cohort study<sup>33</sup> explored the effects of various QI strategies on racial health care disparities in coronary artery disease. The RCT addressed reduction of risk factors for coronary artery disease,<sup>22</sup> while the retrospective cohort examined management of acute myocardial infarction (AMI).<sup>33</sup> QI strategies included patient

education and facilitation of self-management<sup>22</sup> and a multifactorial provider– and systems–focused strategy.<sup>33</sup> Both studies were collaborations of academic and community health centers.<sup>22,33</sup> The studies each employed an internal "usual care" comparison group.

One study of cardiovascular risk factor modification showed no meaningful reduction in health disparities seen in smoking rates, although both Black and White participants had substantially lower rates of smoking after intervention. The other study of AMI treatment reduced disparities in one aspect of treatment, which exacerbated disparities in other areas. The strength of evidence was insufficient.

Two post hoc analyses of RCTs explored the effects of various QI strategies on racial health care disparities in hypertension. The RCTs addressed management of hypertension and reduction of risk factors for coronary artery disease (CAD) including hypertension. QI strategies were patient education and facilitation of self-management. The studies took place in university clinics and multicenter collaborations of academic and community health centers. The studies each employed an internal "usual care" comparison group.

One study had no significant intervention effect on a clinically insignificant disparity in blood pressure measures present at baseline after patient education and promotion of self management. <sup>22</sup> In the second study, however, a home-based self-management strategy, including home blood pressure monitoring and tailored self-management strategies were more effective in the Black population than in the White population, although the study design precludes a clear causal effect by the intervention. <sup>21</sup>

## **Applicability**

Studies of CAD risk factor control included men with CAD risk factors at clinical centers in 18 U.S. cities, and patients with hypertension cared for at two university-affiliated clinics in North Carolina. A study involving treatment of acute myocardial infarction (AMI) included patients hospitalized at academic and community hospitals in Michigan. The study involving men only has limited applicability to women, as patterns of CAD risk factors differ by sex. Moreover, its enrollment occurred between 1973 and 1975, limiting applicability to present day practice. The results of the other two studies are applicable to patients in academic primary care practices, or academic or community hospitals, respectively.

The interventions for CAD risk factor control included intensive patient education and self-management, along with medication titration in one study. The intervention for AMI treatment involved provider education, practice feedback, and implementation of a toolkit. These all required significant institutional resources and the CAD risk factor interventions in particular may not be feasible in routine clinical practice. The AMI treatment initiative, though requiring institutional commitment, has already been disseminated extensively around the United States as a professional society initiative (American College of Cardiology Guidelines Applied in Practice), and thus, its replication is confirmed to be feasible. In each of these studies, usual care served as the comparator. As this varies across practice settings, the effect of the interventions may differ in other environments.

For studies of cardiovascular risk factor control, outcomes consisted of intermediate clinical variables (hypertension, cholesterol, smoking, weight). Outcome assessment in the AMI treatment study was extensive but focused on measures of process and proximal utilization (e.g., prescription of evidence-based medications, use of cardiac catheterization).

# **Depression**

# **Summary**

The three studies that evaluated the effect of QI interventions on disparities in depression outcomes focused on racial disparities but included analyses on sex,<sup>30</sup> income,<sup>18</sup> and educational status.<sup>24</sup> All three studies used a collaborative care model, which involved collaboration among multiple clinical providers to provide a coordinated set of interventions. The clinical model in all three studies included a dedicated mental health coordinator (nurse or case manager), creation of mental health teams (composed of primary provider, facility nurses, and psychiatrists), evidence-based pharmacotherapy and psychotherapy, extensive provider education, and longitudinal patient followup to evaluate clinical status and adherence. Each intervention was designed to address known barriers to the receipt of quality mental health care.

The collaborative care models were all associated with improvements in mental health outcomes, including depression scores, severity and functioning, but none specifically demonstrated a reduction in disparity caused by the intervention. In part, this was because few disparities were measurable at baseline, and what the studies did show is that there was no significant difference in the effect in groups defined by income, race or education. Nonetheless, there were some notable differences in effectiveness that might inform future research. For example, the Prevention of Suicide in Primary Care Elderly: Collaborative Trial (PROSPECT) intervention had a greater effect on clinical outcomes in the less educated group, <sup>24</sup> and the effect of the Partners in Care (PIC) intervention was amplified in minorities on some measures. <sup>28</sup> Although no change in disparity was associated with the interventions, improvements did occur across the board, and no harms were reported in any of the studies. However, because we selected only those studies that could have demonstrated a change in disparity, this review does not include even a small proportion of the overall literature on collaborative care, so it is possible that these studies are anomalies in terms of overall effectiveness.

# **Applicability**

Two of the three studies focused on elderly patients in primary care. One included a range of ages in adulthood. All included both men and women and were racially diverse. Nonetheless, these patient groups may represent a small proportion of the individuals who struggle with depression because of the limited range of health care settings represented in these studies. It is unclear whether the observed results apply to patient populations who receive their primary and mental health care outside of a managed care system, or to individuals who do not receive regular medical care. Additionally, given the settings in which the study took place, they also may not apply to vulnerable populations receiving care through public health systems.

The interventions were all intensive in terms of demand on resources and required strong communication between care providers. In the PIC study, enrolled practiced committed to an intervention cost-sharing arrangement, with the understanding that the long-term implementation would fall on the organization of practice itself. The degree to which this is likely to be feasible is unclear.

All studies compared the intervention to usual care, although usual care was not ever completely described and therefore would be expected to vary.

Generally speaking, outcomes were appropriate and reflected those that would and could be used in practice. They included changes in depressive symptoms, incidence of probably

depressive disorder, mental health related quality of life, functional impairment, and receipt of appropriate depression care.

All studies were conducted in primary care practices associated with larger health care organizations. It is unclear whether results would therefore apply to other settings, including individual practices without the resources of a larger organization, or assisted living facilities (pertinent because of the focus on the elderly population).

#### **Diabetes**

# Summary

Three good quality studies could be used to assess the impact of QI interventions on disparities in diabetes outcomes. One was a randomized controlled trial (RCT),<sup>16</sup> one was a prospective cohort,<sup>31</sup> and one was a retrospective cohort study.<sup>32</sup> None reported critically important clinical outcomes of diabetes, such as death, hypoglycemic coma, adverse drug event, cardiovascular complications, retinopathy progression, nephropathy progression, neurologic complications, or hospitalization for a complication of diabetes. Rather, they reported on surrogate clinical outcomes, clinical risk factors for diabetes comorbidities, and process measures. In two of three studies of diabetes, disparities were reduced in one or more outcomes over the course of the study in at least one subgroup, but the study designs were such that the reduction could not be be shown to be caused by the intervention. <sup>31,32</sup>

In one study of a patient reminder system, racial disparities were reduced when HbA1c testing increased substantially among Black participants, relative to no change among White participants. In a broad, systems level program in New York State, a disparity of 19 percent in biennial lipid testing between African-American beneficiaries and White Medicare recipients was reduced to 9.2 percent after intervention of multifaceted QI program. However, the intervention was multifaceted and widespread, and the authors note that they cannot attribute the change to any specific components of the intervention.

# **Applicability**

Studies included people cared for by primary care clinicians in ambulatory health centers in eastern Massachusetts, diabetes disease management program members living in socioeconomically disparate areas throughout the United States and Medicare patients in New York State. Therefore, the results may or may not be applicable to other populations in other regions.

Interventions evaluated included cultural competency training for clinicians and race-stratified performance reports with recommendations for Black diabetic patients, patient telephone reminders in Health Disparity Zones, defined as one with diabetes prevalence above the national average for minorities, and Medicare New York State Quality Improvement Organization (IPRO) multifaceted provider and community interventions. The interventions may not be available in other regions and settings, since they required significant programmatic and implementation resources. The usual care comparators described in these studies may not be applicable to other settings and regions.

Studies reported surrogate clinical outcomes (i.e., HbA1c control), clinical risk factors for diabetes comorbidities (i.e., blood pressure and lipid control), and process measures (i.e., HbA1c and low density lipoprotein measurements). Duration of studies was generally 1 year. No studies reported any critically important clinical outcomes of diabetes such as death or microvascular

and/or macrovascular complications. Results from surrogate outcomes may not apply to important long-term clinical outcomes in people with diabetes.

Studies were conducted in ambulatory health centers in eastern Massachusetts, in diabetes disease management programs across the United States, and in New York State. As much of diabetes care is delivered in primary care ambulatory settings, the evidence would be applicable. However, specialty clinic settings were not reported and the evidence may not apply.

# **Gaps in the Literature**

A sufficient body of methodologically appropriate research to assess the effects of QI intervention on disparity outcomes currently is not available in the literature. Our assessment is consistent with at least one prior review from 2006, despite several additional years of QI research. Although researchers have focused on the role of QI for improving care in specific populations, there is a fundamental lack of research focused specifically on reducing gaps in the availability, accessibility, and quality of health care between any two populations. Authors of studies in this review have attempted to address the question by conducting post hoc analyses of RCTs intended to study the effectiveness of QI interventions; however, in doing so they have broken what randomization existed and have been unable to make the necessary comparison to tie observed improvements to the intervention conclusively.

Our review specifically sought studies that could measure a potential change in disparity. Other types of studies, such as those that only included only underserved individuals, might provide valuable information to policymakers and clinicians hoping to improve care in those populations, but would not have been included in this review.

## Limitations

One of the challenges to studying the degree to which QI interventions can be found to address disparities is the substantial breadth and heterogeneity of clinical conditions of interest, populations of people with the clinical conditions, QI intervention strategies, comparator, important clinical outcomes, surrogate outcomes, and disparities of interest. Compounding this heterogeneity is poor indexing of QI strategies in the medical literature databases. For example, the subject term "Quality Improvement" was only added to PubMed in 2011; before this time, myriad subject terms were employed to describe the various strategies employed in the QI literature, understandably leading to tremendous variability in how similar studies are categorized in the database. This is partially due to a lack of consistency or agreement on what constitutes a QI intervention – what is available in the literature is often not clearly identified as such and may be multifaceted and thus difficult to evaluate or compare to other intervention studies. Many QI interventions also include non-QI components, such as broader public health initiatives; thus, the potential impact of the QI intervention may be masked or difficult to isolate.

A further challenge to studying changes in disparities is the poor documentation of those disparities, and the fact that any two populations may represent multiple and overlapping disparities. In the studies that we were able to find that could have empirically assessed a disparity change, many were unable to demonstrate any existing disparity at baseline, which may be a reflection of the complexity of identifying and measuring disparities using single characteristics. Future studies will require much broader populations to include enough individuals from diverse background to capture and assess disparities over time empirically.

Finally, the degree to which publication bias may exist is unknown but potentially important for this literature. Many QI interventions are programmatic interventions in health care systems,

not necessarily designed as research. Decisions about whether to publish the results of such interventions could, therefore, be based on the degree to which they were perceived to be successful and potentially useful to other industry colleagues and systems. It is possible that other interventions have been performed and remain unpublished.

### **Future Research**

Calls for the study of QI strategies to address disparities are not new. In order to advance this field, research must be specifically designed to assess the relevant contrasts in disparities and not rely on post hoc analyses of interventions designed to improve the health and health care of all participating individuals. In part, this is complicated by a lack of detailed evidence about the root causes of disparities.

To determine if a QI intervention is effective for reduction of a disparity, the research protocol needs to establish the baseline health outcomes for the health condition of interest for at least two groups, and this baseline data must include the report of disparity. The research protocol must be designed to produce a report of the same health outcome measures after the QI intervention for the same groups, and that post-intervention analysis must include data about the disparity. In this way, two levels of effectiveness are measured simultaneously—that for the intervention overall and that for the reduction of disparity.

For intervention effectiveness to be established there must be a statistically significant improvement in the health outcomes for an intervention group, compared with a control group. For reduction of disparity effectiveness to be established, there must be a statistically significant reduction of the disparity measured after the intervention, compared with the baseline disparity; complex sample size calculations must be performed to determine that the study will be adequately powered for both QI intervention effectiveness versus control, and for QI intervention effectiveness for reduction of disparity. A potentially effective intervention for reducing inequities may be one that is equally effective across the socioeconomic spectrum, but that may reduce health inequalities simply because the prevalence of health problems among the disadvantaged is greater. It may also be the case that an intervention is more effective in a disparity group because it addresses their particular needs, but also benefits the non-disparity group (e.g., a disease management program that pays attention to issues of health literacy may provide greater benefits to patients with low health literacy). Nonetheless, the vast majority of studies purported to address disparities study only one group—the disadvantaged group—thus eliminating the studies' own ability to actually measure an actual change in disparity.

Even with well-designed studies, the basis for believing that a QI intervention designed to improve health care overall would reduce a disparity is the assumption that it would have an amplified effect in a disadvantaged group, thus accelerating their health care improvement to the point that the gap decreases. This assumption has not been adequately studied in any health care condition or in relation to any disparity, and this foundational research is necessary to establish the basis for continuing research on QI interventions with the expectation that they will affect disparities.

Nonetheless, QI interventions that demonstrate improvements across all populations do not necessarily do so at the expense of disadvantaged groups; they simply do not demonstrate an ability to accelerate change in one group versus another to the point that a gap is closed. Therefore, it is unclear whether QI interventions are potentially fruitful or appropriate for the purpose of reducing health care outcome disparities, despite their known effectiveness at improving health and health care across disparity groups.

Nonetheless, research should be conducted on the collaborative care model and the role of targeted patient education, which is accommodating of recipients' language and literacy levels. As previously noted, sufficient data are not available to support universal implementation of these strategies, but the strategies may be suitable for implementation if an appropriate plan is in place to monitor their effectiveness. In particular, it will be important in studying these and other QI interventions to ensure capture on data reflecting organizational adherence to interventions, something notably missing from the current literature.

Furthermore, data on patient characteristics known to be associated with disparity should be collected regularly such that any observed changes in disparity can be measured. Future studies should provide data in such a way that both the overall effectiveness and a change in disparity can be assessed statistically.

## **Conclusions**

The literature on QI interventions generally and their ability to improve health and health care is large. Whether those interventions are effective at reducing disparities remains unclear. This report should not be construed to assess the general effectiveness of QI in the health care setting; rather, QI has not been shown specifically to reduce known disparities in health care or health outcomes. In a few instances, some increased effect is seen in disadvantaged populations; these studies should be replicated and the interventions studied further as having potential to address disparities. Specific examples that warrant additional study include increasing our understanding of the collaborative care model in improving depression outcomes, and the role of targeted patient education, including ensuring that interventions are provided in the primary language of the patients.

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# **Acronyms and Abbreviations**

AA African-American

ADA American Diabetes Association

AHRQ Agency for Healthcare Research and Quality

AMI Acute myocardial infarction

BP Blood pressure

CABG Coronary artery bypass grafting

CAD Coronary artery disease CBT Cognitive behavior therapy

CC Collaborative care
CHD Coronary heart disease
CI Confidence interval
CQG Closing the Quality Gap

CQI Continuous Quality Improvement

CRC Colorectal cancer

DBP Diastolic blood pressure
DCM Depression care manager
DCS Depression care specialist
EPC Evidence-based practice center

FADE Quality improvement organizational model: Focus, Analyze, Develop, Execute,

**Evaluate** 

FFS Fee-for-service

FOBT Fecal occult blood test

FS/COL Flexible sigmoidoscopy or colonoscopy

GAP American College of Cardiology's AMI Guidelines Applied in Practice

GCC Guideline concordant care
HbA1c Glycosylated hemoglobin
HDL High density lipoprotein
HDZ Health disparity zone

HMO Health maintenance organization

IMPACT Improving mood-promoting access to collaborative treatment

IOM Institute of Medicine

IPRO Medicare quality improvement organization for the State of New York

KQ Key question

LDL Low density lipoprotein

MCO Managed care organization

MCS-12 Mental health component scale

MDD Major depressive disorder

mm Hg Millimeter of mercury

mmol/L Millimole per liter

MRFIT Multiple risk factor intervention trial

N-O Newcastle-Ottawa NA Not applicable

Non-HDZ Non-health disparity zone

NP Nurse practitioner

NR Not reported

adjusted OR Adjusted odds ratio

OR Odds ratio

PA Physician assistant

PCI Percutaneous coronary intervention

PCS-12 Physical component score PDSA Plan, Do, Study, Act PIC Partners in care

PROSPECT Prevention of Suicide in Primary Care Elderly

QI Quality improvement

QI–Meds Quality improvement-enhanced medication management QI–Therapy Quality improvement-enhanced therapy management

QoL Quality of life

RCT Randomized controlled trial

REALM Rapid estimate of adult literacy in medicine

RoB Risk of bias

SBP Systolic blood pressure
SD Standard deviation
SES Socioeconomic status
TEP Technical Expert Panel
TQM Total quality management

UC Usual care VA Veterans Affairs

# **Appendix A. Literature Search Strategies**

Database: PubMed

Search	Search terms	Search results
#1	reminder systems[mh] OR guideline adherence[mh] OR medical audit[mh] OR interdisciplinary communication[mh] OR feedback[mh] OR nursing audit[mh] OR patient education as topic[mh] OR education, continuing[mh] OR health personnel/education[mh] OR health education[majr] OR "provider education" OR self care[mh] OR organizational innovation[mh] OR "self management" OR quality improvement[tiab] OR quality assurance, health care[mh] OR quality indicators, health care[mh] OR safety management[majr] OR patient safety[tiab]	481284
#2	health status disparities[mh] OR healthcare disparities[mh] OR minority health[mh] OR ethnic groups[mh] OR minority groups[mh] OR health literacy[mh] OR health literacy[tiab] OR numeracy[tiab] OR socioeconomic factors[mh] OR social class[mh] OR sexuality[mh] OR communication barriers[mh] OR translating[mh] OR language[mh:noexp] OR insurance coverage[mh] OR medically uninsured[mh] OR disparities[tiab] OR disparity[tiab] OR inequality[tiab] OR inequalities[tiab]	475923
#3	colorectal neoplasms[mh] OR breast neoplasms[mh] OR heart failure[mh] OR myocardial ischemia[mh] OR coronary disease[mh] OR diabetes mellitus[mh:noexp] OR diabetes mellitus, type 1[mh] OR diabetes mellitus, type 2[mh] OR hypertension[mh:noexp] OR pregnancy outcome[mh] OR birth weight[mh] OR premature birth[mh] OR infant mortality[mh] OR infant, low birth weight[mh] OR infant, premature[mh] OR depressive disorder[mh:noexp] OR depressive disorder, major[mh] OR depression[mh] OR asthma[mh] OR pneumonia[mh] OR pneumococcal vaccines[mh] OR cystic fibrosis[mh] OR kidney failure, chronic[mh] OR renal dialysis[mh]	1517358
#4	#1 AND #2 AND #3 AND eng[la] AND humans[mh] AND 1983:2011[dp]	3616
#5	#4 AND case reports[pt]	44
#6	#4 AND letter[pt]	49
#7	#4 AND review[pt]	288
#8	#4 AND editorial[pt]	43
#9	#4 AND comment[pt]	74
#10	#4 AND practice guideline[pt]	16
#11	#4 AND historical article[pt]	11
#12	#4 AND legal cases[pt]	2
#13	#4 AND news[pt]	27
#14	#4 AND newspaper article[pt]	2
#15	#4 AND meta-analysis[pt]	15
#16	#4 AND jsubsetk	16
#17	#5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16	507
#18	#4 NOT #17	3109

**Key:** jsubsetk consumer health subset; [la] language; [mh] Medical Subject Heading; [mh:noexp] Medical Subject Heading, not including narrower terms; [pt] publication type; [tiab] title or abstract word

# **Database: CINAHL (EBSCOhost interface)**

Search	Search terms	Search results
#1	(MH "Quality Assurance") OR (MH "Quality of Health Care") OR (MH "Patient Safety") OR (MH "Reminder Systems") OR (MH "Health Education+") OR (MH "Audit") OR (MH "Feedback") OR (MH "Self Care") OR TX quality OR (MH "Education, Continuing") OR (MH "Education, Medical, Continuing") OR (MH "Education, Nursing, Continuing") OR (MH "Quality Management, Organizational") OR (MH "Organizational Change")	271475
#2	(TX disparity OR disparities OR numeracy OR health literacy OR inequity OR inequities OR inequality OR inequalities) OR (MH "Minority Groups") OR (MH "Socioeconomic Factors+") OR (MH "Communication Barriers") OR (MH "English as a Second Language") OR (MH "Insurance, Health+") OR (MH "GLBT Persons+") OR (MH "Sexuality") OR (MH "Bisexuality") OR (MH "Homosexuality") OR (MH "Medically Uninsured") OR (MH "Ethnic Groups+")	241818
#3	(MH "colorectal neoplasms+") OR (MH "breast neoplasms+") OR (MH "heart failure") OR (MH "myocardial ischemia+") OR (MH "diabetes mellitus, insulin-dependent") OR (MH "diabetes mellitus, non-insulin-dependent") OR (MH "hypertension") OR (MH "infant, low birth weight") OR (MH "infant, premature") OR (MH "pregnancy outcomes") OR (MH "infant mortality") OR (MH "depression") OR (MH "asthma") OR (MH "pneumonia+") OR (MH "pneumococcal vaccine") OR (MH "cystic fibrosis") OR (MH "kidney failure, chronic") OR (MH "hemodialysis")(MH "colorectal neoplasms+") OR (MH "breast neoplasms+") OR (MH "heart failure") OR (MH "myocardial ischemia+") OR (MH "diabetes mellitus, insulin-dependent") OR (MH "diabetes mellitus, non-insulin-dependent") OR (MH "hypertension") OR (MH "infant, low birth weight") OR (MH "infant, premature") OR (MH "pregnancy outcomes") OR (MH "infant mortality") OR (MH "depression") OR (MH "asthma") OR (MH "pneumonia+") OR (MH "pneumococcal vaccine") OR (MH "cystic fibrosis") OR (MH "kidney failure, chronic"	208909
#4	#1 AND #2 AND #3, limited to English, human, peer reviewed journals, research articles; MEDLINE articles excluded	413
#5	#4 AND (PT "abstract")	55
#6	#4 AND (MH "case studies")	10
#7	#4 AND (PT "commentary")	3
#8	#4 AND (PT "editorial")	1
#9	#4 AND (PT "letter")	1
#10	#4 AND (PT "practice guidelines")	1
#11	#4 AND (PT "proceedings")	1
#12	#5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11	70
#13	#4 NOT #12	343

Key: MH CINAHL medical subject heading; + explode term; PT publication type; TX text word

## **Database: PsycINFO (CSA interface)**

Search	Search terms	Search results
#1	DE=(quality of care OR clinical audits OR feedback OR knowledge of results OR client education OR continuing education OR decision support systems OR self care skills OR self management OR organizational innovation OR quality control OR safety OR "health education") OR (reminder OR reminders), limited to 1980 -2011, journal article, English language, and human	35168
#2	DE=(health disparities OR racial and ethnic groups OR racial and ethnic differences OR health literacy OR minority groups OR english as a second language OR communication barriers OR language proficiency OR numerical ability OR "health insurance" or "employee health insurance" or "workers compensation insurance" or "fee for service" or "health maintenance organizations" or "medicaid" or "medicare" or "underinsured health insurance" or "uninsured health insurance" OR "transgender" or "transsexualism" OR "sexual orientation" or "bisexuality" or "heterosexuality" or "homosexuality" or "lesbianism" or "male homosexuality") OR (disparity OR disparities OR inequity OR inequities OR inequality OR inequalities), limited to 1980 -2011, journal article, English language, and human	80091
#3	DE=(breast neoplasms OR hypertension OR pregnancy outcomes OR premature birth OR birth weight OR asthma OR pneumonia OR cystic fibrosis OR kidney diseases OR "heart disorders" or "angina pectoris" or "arrhythmias heart" or "bradycardia" or "fibrillation heart" or "tachycardia" or "coronary thromboses" or "myocardial infarctions" OR "diabetes" or "diabetes mellitus" OR "major depression" OR "depression emotion" OR "dialysis" or "hemodialysis") OR (DE=(neoplasms) AND (DE=(colon disorders) OR colon OR colorectal)) OR (pneumococcal AND (vaccine OR vaccines OR vaccination OR vaccinated OR vaccinations)), limited to 1980 - 2011, journal article, English language, and human	80009
#4	#1 AND #2 AND #3	391
#5	PT=(edited book) or PT=(editorial) or PT=(electronic collection) or PT=(encyclopedia entry) or PT=(encyclopedia) or PT=(erratum/correction) or PT=(handbook/manual) or PT=(letter) or PT=(obituary) or PT=(publication information) or PT=(reference book) or PT=(reprint) or PT=(review-book) or PT=(review-media) or PT=(textbook/study guide) or PT=(dissertation) or PT=(review-software) or PT=(abstract collection) or PT=(authored book) or PT=(bibliography) or PT=(book) or PT=(chapter) or PT=(classic book) or PT=(column/opinion) or PT=(comment/reply) or PT=(conference proceedings) or PT=(dissertation abstract)	1101
#6	ME=(focus group OR interview OR literature review OR meta analysis OR nonclinical case study OR systematic review)	94944
#7	#4 NOT (#5 OR #6)	336

**Key:** DE subject term; PT publication type; ME methodology

## **Database: Social Science Citation**

Search	Search terms	Search results
#1	TS=(health literacy OR (language AND (barrier* OR problem*)) OR English as a second language OR (health AND numeracy)) AND Language=(English) AND Document Type=(Article)	11111
#2	TS=(breast cancer OR colorectal cancer OR colon cancer OR diabetes OR heart failure OR coronary artery disease OR heart disease OR myocardial infarction OR myocardial ischemia OR hypertension OR hypertensive OR depression OR depressive disorder OR asthma OR cystic fibrosis OR pneumonia OR end stage renal disease OR (birthweight AND low) OR premature birth OR prematurity OR preterm birth) AND Language=(English) AND Document Type=(Article)	>100000
#3	TS=(reminder OR reminders OR audit OR feedback OR patient education OR provider education OR self management OR quality OR patient safety) AND Language=(English) AND Document Type=(Article)	>100000
#4	#1 AND #2 AND #3 , limited to articles with any US author affiliation	280

Key: TS keyword

# **Appendix B. Abstract Review Form**

Please complete each item below (items 1-6) irrespective of the response to the previous item.

Pri	Primary Inclusion/Exclusion Criteria					
1.	Original research (exclude reviews, systematic reviews, editorials, commentaries, letters to editor, etc.).	Yes	No	Cannot Determine		
2.	Includes an intervention.	Yes	No	Cannot Determine		
3.	Includes individuals receiving health care within the U.S.	Yes	No	Cannot Determine		
4.	The number of participants enrolled is greater than or equal to 50 per group.	Yes	No	Cannot Determine		
5.	Addresses one or more of the priority conditions (check one or more):  a Colorectal cancer including screening b Breast cancer including screening c Diabetes mellitus d Congestive heart failure (i.e. heart failure, left-sided heart failure, right-sided heart failure, cor pulmonale, CHF) e Coronary artery disease (i.e. coronary heart disease, arteriosclerotic heart disease, CHD, CAD) f Hypertension g Pregnancy h Major depressive disorder i Asthma j Cystic fibrosis k Pneumonia including pneumococcal vaccination l End stage renal disease	Yes	No	Cannot Determine		
6.	Do the participants include individuals from a target population defined by one of the following indicators of disparity:  a Race/ethnicity b Socioeconomic status c Insurance status d Sex e Sexual orientation f Health literacy/numeracy g Language barrier	Yes	No	Cannot Determine		

If "No" was marked for any response above, the form is complete\*; otherwise, turn the sheet over and complete Part II

*If not included, the citation may still be marked for retention for one of the following reasons:
BACKGROUND/DISCUSSION
REVIEW OF REFERENCES
OTHER

## Please complete each item below (items 7-10) irrespective of the response to the previous item.

7.	Addresses one or more quality improvement strategies (i.e. a systematic process designed to improve the quality of care) as an intervention (check one or more):  a Provider reminder systems b Facilitated relay of clinical data to providers c Audit and feedback d Provider education e Patient education f Promotion of self-management g Patient reminder systems h Other (list below, excluding financial incentives and public reporting):	Yes	No	Cannot Determine
8.	Intervention originates from or occurs within at least one of the following settings:  a Hospital  b Clinic  c Provider office	Yes	No	Cannot Determine
9.	Includes outcomes of interest for a referent group that is either an:  a Internal source (i.e. within study referent group) or b External source (i.e. data from a referent group not included in the study)	Yes	No	Cannot Determine
10.	Addresses an outcome of interest (check one or both):			
	a Health outcome and/or process outcome b Harm and/or unanticipated adverse effect	Yes	No	Cannot Determine
	If "No" was marked for any of the items (#7-#10) above, and the citation is excluded by review, the citation will not proceed to full text review.*  *If not included, the citation may still be marked for retention for one of the following remarked.  BACKGROUND/DISCUSSION  REVIEW OF REFERENCES  OTHER		abstract	

# **Appendix C. Full-Text Review Form**

Please complete each item below (items 1-4) irrespective of the response to the previous item.

1.	Does the paper describe original research (i.e. the paper is not a review article, meta-analysis, systematic review, editorial, commentary, letter to the editor, patient summary, etc.)?			No
2.	Does the intervention or strategy meet the definition of quality improvement?			
	If "yes", check the definition that applies:			
	Quality improvement definition		Yes	No
	$\hfill\Box$ A formal, broad organization model, such as PDSA (plaimprovement), or TQM (total quality management).	an, do, study, act), Six Sigma, CQI (continuous quality	165	INO
	$\hfill \square$ A change process in health care systems, services, or soptimal clinical quality of care.	supplier for the purpose of increasing the likelihood of		
	Select one or more of the QI categories below.			
	Quality improvement taxonomy			
	<ul> <li>□ Provider reminder system</li> <li>□ Facilitated relay of clinical data to provider</li> <li>□ Audit and feedback</li> <li>□ Provider education</li> </ul>	<ul> <li>□ Patient education/Promotion of self-management</li> <li>□ Patient reminder system</li> <li>□ Other</li> </ul>		
3.	Does the study target disparities that are based on one of	the selected indicators?		
	If "yes", check all that apply below:		Yes	No
	Disparity indicator(s)			1
	☐ Race/ethnicity	☐ Sexual orientation		
	☐ Socioeconomic status	☐ Health literacy/numeracy		
	☐ Insurance status	☐ Language barrier		
	□ Sex			
	Does the study demonstrate or measure a change in disparenral) comprising individuals without the disparity of interes			
	If "yes", check one of the data sources below:			
	Referent data source		Yes	No
	$\hfill\Box$ Internal (i.e. a group of individuals within the study)			
	$\hfill\Box$ External (i.e. data from a source or group outside the st	udy)		
4ai. If the referent data source is external, does it meet both the following conditions?				
a) geographically local (i.e. not greater than state level); and b) temporally proximal (i.e. not greater than four years from the date of target group data collection)			Yes	No
	Does the study report data from the referent group (internal quality improvement intervention?	or external) both before and after the introduction of	Yes	No

If "no" is checked for one or more of the items above, the form is complete. Otherwise, continue and complete items 5-7 below.

		1					
5. Does the study meet the specified conditions (see below) for inclusion?							
If "no", check one or more of the conditions <b>NOT</b> met below:							
Exclusion reason							
☐ Study participants or centers are not based in the U.S.		Yes	No				
☐ The intervention does not originate from a clinic, hospit	tal, or provider office.						
☐ The study does not include a control group for the qual	lity improvement strategy/intervention.						
☐ The study does not include 50 or more participants for	each intervention and control group.						
6. Does the study address a priority condition?							
If "yes", check one or more of the conditions below:		Yes	No				
Condition(s)							
☐ Colorectal cancer (including screening)							
☐ Breast cancer (including screening) ☐ Pregnancy (excluding neonatal only)							
☐ Diabetes mellitus (insulin and non-insulin dependent)							
☐ Congestive heart failure							
☐ Cystic fibrosis							
disease, acute coronary syndrome, acute myocardial	☐ Pneumonia (including pneumococcal vaccination)						
infarction, and STEMI)							
☐ Hypertension							
7. Does the study address an outcome(s) of interest?							
If "yes", check one or more outcome categories listed below:							
Outcome category		Yes	No				
☐ Health outcome and/or process outcome							
☐ Harm and/or unanticipated adverse effect							
<u>Optional</u>							
Should this article be retained for background information, review of r If "yes", check one or more reasons below.	references, or other reasons?						
Retain for:							
$\square$ BACKGROUND/DISCUSSION $\square$ REVIEW OF REFEREN	ICES  OTHER						
<u>Optional</u>							
Enter comments or notes below.							

C-2

# **Appendix D. Cochrane Risk of Bias Tool**

Cochrane Collaboration modified tool for assessing risk of bias for RCT's, PART I Use this form to assess risk of bias for randomized controlled trials.

Bias is assessed as a judgement (high, low, or unclear) for individual elements from five domains (selection, performance, attrition, reporting, and other).

Risk of selection, reporting, and other bias are assessed in the **Quality Assessment Form Part I.** Risk of performance, detection, and attrition bias are assessed using the **Quality Assessment Form Part II.** 

Using the guidance provided at the end of this form, select either "high", "low" or "unclear" for each judgment. When complete, proceed to **Part II of the Quality Assessment Form** 

REF ID:					
Domain	Description	High risk of bias	Low risk of bias	Unclear risk of bias	Reviewer Assessment
Selection bias Random sequence generation	Described the method used to generate the allocation sequence in sufficient detail to allow an assessment of whether it should produce comparable groups.  Reviewer Comments:	Selection bias (biased allocation to interventions) due to inadequate generation of a randomized sequence.	Random sequence generation method should produce comparable groups	Not described in sufficient detail	Judgement  Random sequence generation  High Low Unclear
Selection bias Allocation concealment	Described the method used to conceal the allocation sequence in sufficient detail to determine whether intervention allocations could have been foreseen in advance of, or during, enrollment.  Reviewer Comments:	Selection bias (biased allocation to interventions) due to inadequate concealment of allocations prior to assignment.	Intervention allocations likely could not have been foreseen in advance of, or during, enrollment	Not described in sufficient detail	Judgement Allocation concealment  High Low Unclear
Reporting bias  Selective reporting	Stated how the possibility of selective outcome reporting was examined by the authors and what was found.  Reviewer Comments:	Reporting bias due to selective outcome reporting.	Selective outcome reporting bias not detected	Insufficient information to permit judgement (It is likely that the majority of studies will fall into this category.)	Judgement Selective reporting  High Low Unclear
Other bias  Other sources of bias	Any important concerns about bias not addressed above. If particular questions/entries were pre-specified in the study's protocol, responses should be provided for each question/entry.  Reviewer Comments:	Bias due to problems not covered elsewhere in the table.	No other bias detected	There may be a risk of bias, but there is either insufficient information to assess whether an important risk of bias exists; or insufficient rationale or evidence that an identified problem will introduce bias.	Judgement Other sources of bias  High Low Unclear

#### Cochrane Collaboration modified tool for assessing risk of bias for RCT's, PART II

Use this form to assess risk of bias for randomized controlled trials.

Bias is assessed as a judgement (high, low, or unclear) for individual elements from five domains of bias (selection, performance, attrition, reporting, and other).

Risk of selection, reporting, and other bias are assessed in the **Quality Assessment Form Part I.** Risk of performance, detection, and attrition bias are assessed using the **Quality Assessment Form Part II.** 

Using the guidance provided at the end of this form, select either "high", "low" or "unclear" for each judgement.

Risk of bias for the domains in the Form Part II will be assessed for each main or class of outcomes. Please indicate the specific outcome and complete the assessment for each.

REF ID:					
Outcomes:					
Domain	Description	High risk of bias	Low risk of bias	Unclear risk of bias	Reviewer Assessment
Performance bias Blinding (participants and personnel)	Described all measures used, if any, to blind study participants and personnel from knowledge of which intervention a participant received. Provided any information relating to whether the intended blinding was effective.  Reviewer Comments:	Performance bias due to knowledge of the allocated interventions by participants and personnel during the study.		Not described in sufficient detail	Judgement  Blinding (participants and personnel)
Detection bias  Blinding (outcome assessment)	Described all measures used, if any, to blind outcome assessors from knowledge of which intervention a participant received. Provided any information relating to whether the intended blinding was effective.  Reviewer Comments:	Detection bias due to knowledge of the allocated interventions by outcome assessors.	Blinding was likely effective.	Not described in sufficient detail	Judgement  Blinding (outcome assessment)
Attrition bias Incomplete outcome data	Described the completeness of outcome data for each main outcome, including attrition and exclusions from the analysis. Stated whether attrition and exclusions were reported, the numbers in each intervention group (compared with total randomized participants), reasons for attrition/exclusions where reported.  Reviewer Comments:	Attrition bias due to amount, nature or handling of incomplete outcome data.	incomplete outcome data was complete and unlikely to	Insufficient reporting of attrition/exclusions to permit judgment of 'Low risk' or 'High risk' (e.g. number randomized not stated, no reasons for missing data provided)	Judgement Incomplete outcome data

# **Appendix E. Cochrane Risk of Bias Criteria**

Criteria for judging risk of bias in the 'Risk of bias' assessment tool\*

Bias	Judgment	Criteria
RANDOM	'Low risk' of bias.	The investigators describe a random component in the sequence generation process such as:  Referring to a random number table; Using a computer random number generator; Coin tossing; Shuffling cards or envelopes; Throwing dice; Drawing of lots; Minimization*.  *Minimization may be implemented without a random element, and this is considered to be equivalent to being random.
SEQUENCE GENERATION Selection bias (biased allocation to interventions) due to inadequate generation of a randomised sequence.	'High risk' of bias.	The investigators describe a non-random component in the sequence generation process. Usually, the description would involve some systematic, non-random approach, for example:  • Sequence generated by odd or even date of birth;  • Sequence generated by some rule based on date (or day) of admission;  • Sequence generated by some rule based on hospital or clinic record number.  Other non-random approaches happen much less frequently than the systematic approaches mentioned above and tend to be obvious. They usually involve judgement or some method of non-random categorization of participants, for example:  • Allocation by judgement of the clinician;  • Allocation by preference of the participant;  • Allocation based on the results of a laboratory test or a series of tests;  • Allocation by availability of the intervention.
	'Unclear risk' of bias.	Insufficient information about the sequence generation process to permit judgement of 'Low risk' or 'High risk'.
ALLOCATION CONCEALMENT Selection bias	'Low risk' of bias.	Participants and investigators enrolling participants could not foresee assignment because one of the following, or an equivalent method, was used to conceal allocation:  • Central allocation (including telephone, web-based and pharmacy-controlled randomization);  • Sequentially numbered drug containers of identical appearance;  • Sequentially numbered, opaque, sealed envelopes.
(biased allocation to interventions) due to inadequate concealment of allocations prior to assignment.	'High risk' of bias.	Participants or investigators enrolling participants could possibly foresee assignments and thus introduce selection bias, such as allocation based on:  • Using an open random allocation schedule (e.g. a list of random numbers);  • Assignment envelopes were used without appropriate safeguards (e.g. if envelopes were unsealed or nonopaque or not sequentially numbered);  • Alternation or rotation;  • Date of birth;  • Case record number;  • Any other explicitly unconcealed procedure.

Bias	Judgment	Criteria
	'Unclear risk' of bias.	Insufficient information to permit judgement of 'Low risk' or 'High risk'. This is usually the case if the method of concealment is not described or not described in sufficient detail to allow a definite judgement – for example if the use of assignment envelopes is described, but it remains unclear whether envelopes were sequentially numbered, opaque and sealed.
	'Low risk' of bias.	Any of the following:  The study protocol is available and all of the study's pre-specified (primary and secondary) outcomes that are of interest in the review have been reported in the pre-specified way;  The study protocol is not available but it is clear that the published reports include all expected outcomes, including those that were pre-specified (convincing text of this nature may be uncommon).
SELECTIVE REPORTING Reporting bias due to selective outcome reporting.	'High risk' of bias.	<ul> <li>Any one of the following:</li> <li>Not all of the study's pre-specified primary outcomes have been reported;</li> <li>One or more primary outcomes is reported using measurements, analysis methods or subsets of the data (e.g. subscales) that were not pre-specified;</li> <li>One or more reported primary outcomes were not pre-specified (unless clear justification for their reporting is provided, such as an unexpected adverse effect);</li> <li>One or more outcomes of interest in the review are reported incompletely so that they cannot be entered in a meta-analysis;</li> <li>The study report fails to include results for a key outcome that would be expected to have been reported for such a study.</li> </ul>
	'Unclear risk' of bias.	Insufficient information to permit judgement of 'Low risk' or 'High risk'. It is likely that the majority of studies will fall into this category.
	'Low risk' of bias.	The study appears to be free of other sources of bias.
OTHER BIAS Bias due to problems not covered elsewhere	'High risk' of bias.	There is at least one important risk of bias. For example, the study:  • Had a potential source of bias related to the specific study design used; or  • Has been claimed to have been fraudulent; or  • Had some other problem.
in the table.	'Unclear risk' of bias.	There may be a risk of bias, but there is either:  Insufficient information to assess whether an important risk of bias exists; or  Insufficient rationale or evidence that an identified problem will introduce bias.
BLINDING OF PARTICIPANTS AND PERSONNEL	'Low risk' of bias.	<ul> <li>Any one of the following:</li> <li>No blinding or incomplete blinding, but the review authors judge that the outcome is not likely to be influenced by lack of blinding;</li> <li>Blinding of participants and key study personnel ensured, and unlikely that the blinding could have been broken.</li> </ul>
Performance bias due to knowledge of the allocated interventions by participants and personnel during	'High risk' of bias.	<ul> <li>Any one of the following:</li> <li>No blinding or incomplete blinding, and the outcome is likely to be influenced by lack of blinding;</li> <li>Blinding of key study participants and personnel attempted, but likely that the blinding could have been broken, and the outcome is likely to be influenced by lack of blinding.</li> </ul>
the study.	'Unclear risk' of bias.	Any one of the following:  Insufficient information to permit judgment of 'Low risk' or 'High risk';  The study did not address this outcome.

Bias	Judgment	Criteria
BLINDING OF OUTCOME ASSESSMENT	'Low risk' of bias.	Any one of the following:     No blinding of outcome assessment, but the review authors judge that the outcome measurement is not likely to be influenced by lack of blinding;     Blinding of outcome assessment ensured, and unlikely that the blinding could have been broken.
Detection bias due to knowledge of the allocated interventions by outcome assessors.	'High risk' of bias.	Any one of the following:     No blinding of outcome assessment, and the outcome measurement is likely to be influenced by lack of blinding;     Blinding of outcome assessment, but likely that the blinding could have been broken and the outcome measurement is likely to be influenced by lack of blinding.
	'Unclear risk' of bias.	Any one of the following:  Insufficient information to permit judgment of 'Low risk' or 'High risk';  The study did not address this outcome.
INCOMPLETE	'Low risk' of bias.	<ul> <li>Any one of the following: <ul> <li>No missing outcome data;</li> <li>Reasons for missing outcome data unlikely to be related to true outcome (for survival data, censoring unlikely to be introducing bias);</li> <li>Missing outcome data balanced in numbers across intervention groups, with similar reasons for missing data across groups;</li> <li>For dichotomous outcome data, the proportion of missing outcomes compared with observed event risk not enough to have a clinically relevant impact on the intervention effect estimate;</li> <li>For continuous outcome data, plausible effect size (difference in means or standardized difference in means) among missing outcomes not enough to have a clinically relevant impact on observed effect size;</li> <li>Missing data have been imputed using appropriate methods.</li> </ul> </li> </ul>
OUTCOME DATA Attrition bias due to amount, nature or handling of incomplete outcome data.	'High risk' of bias.	<ul> <li>Any one of the following:</li> <li>Reason for missing outcome data likely to be related to true outcome, with either imbalance in numbers or reasons for missing data across intervention groups;</li> <li>For dichotomous outcome data, the proportion of missing outcomes compared with observed event risk enough to induce clinically relevant bias in intervention effect estimate;</li> <li>For continuous outcome data, plausible effect size (difference in means or standardized difference in means) among missing outcomes enough to induce clinically relevant bias in observed effect size;</li> <li>'As-treated' analysis done with substantial departure of the intervention received from that assigned at randomization;</li> <li>Potentially inappropriate application of simple imputation.</li> </ul>
	'Unclear risk' of bias.	Any one of the following:  Insufficient reporting of attrition/exclusions to permit judgement of 'Low risk' or 'High risk' (e.g. number randomized not stated, no reasons for missing data provided);  The study did not address this outcome.

<sup>\*</sup> Adapted from the Cochrane Collaboration Risk of Bias Criteria

# Appendix F. Newcastle-Ottawa Quality Assessment Scale

Ass	essment of quality of a cohort study – Newcastle-Ottawa Scale						
Sele	Selection (tick one box in each section)						
1.	Representativeness of the intervention cohort  a) truly representative of the <u>average</u> , <u>elderly</u> , <u>community-dwelling resident</u> b) somewhat representative of the <u>average</u> , <u>elderly</u> , <u>community-dwelling resident</u> c) selected group of patients, <u>e.g. only certain socio-economic groups/areas</u> d) no description of the derivation of the cohort						
2.	Selection of the non intervention cohort  a) drawn from the same community as the intervention cohort  b) drawn from a different source  c) no description of the derivation of the non intervention cohort						
3.	Ascertainment of intervention  a) secure record (eg health care record)  b) structured interview  c) written self report  d) other / no description						
4.	Demonstration that outcome of interest was not present at start of study a) yes b) no						
Con	nparability (tick one or both boxes, as appropriate)						
1.	Comparability of cohorts on the basis of the design or analysis  a) study controls for <u>age, sex, marital status</u> b) study controls for any additional factors ( <u>e.g. socio-economic status, education</u> )						
Out	come (tick one box in each section)						
1.	Assessment of outcome  a) independent blind assessment b) record linkage c) self report d) other / no description						
2.	Was follow up long enough for outcomes to occur  a) yes, if median duration of follow-up >= 6 month  b) no, if median duration of follow-up < 6 months						
3.	Adequacy of follow up of cohorts  a) complete follow up: all subjects accounted for  b) subjects lost to follow up unlikely to introduce bias: number lost <= 20%, or description of those lost suggesting no different from those followed  c) follow up rate < 80% (select an adequate %) and no description of those lost d) no statement	0000					

#### NOS - CODING MANUAL FOR COHORT STUDIES

#### **SELECTION**

#### 1) Representativeness of the Exposed Cohort (NB exposure = intervention)

Item is assessing the representativeness of exposed individuals in the community, not the representativeness of the study sample from some general population. For example, subjects derived from groups likely to contain exposed people are likely to be representative of exposed individuals, while they are not representative of all people the community.

Allocation of points as per rating sheet

#### 2) Selection of the Non-Exposed Cohort

Allocation of points as per rating sheet

#### 3) Ascertainment of Exposure

Allocation of points as per rating sheet

#### 4) Demonstration That Outcome of Interest Was Not Present at Start of Study

In the case of mortality studies, outcome of interest is still the presence of a disease/ incident, rather than death. That is to say that a statement of no history of disease or incident earns a point.

#### **COMPARABILITY**

#### 1) Comparability of Cohorts on the Basis of the Design or Analysis

Either exposed and non-exposed individuals must be matched in the design and/or confounders must be adjusted for in the analysis. Statements of no differences between groups or that differences were not statistically significant are not sufficient for establishing comparability. Note: If the relative risk for the exposure of interest is adjusted for the confounders listed, then the groups will be considered to be comparable on each variable used in the adjustment.

A maximum of 2 points can be allotted in this category.

#### **OUTCOME**

#### 2) Assessment of Outcome

For some outcomes, reference to the medical record is sufficient to satisfy the requirement for confirmation. This may not be adequate for other outcomes where reference to specific tests or measures would be required.

- a) Independent or blind assessment stated in the paper, or confirmation of the outcome by reference to secure records (health records, etc.)
- b) Record linkage (e.g. identified through ICD codes on database records)
- c) Self-report (i.e. no reference to original health records or documented source to confirm the outcome)
- d) No description.

#### 3) Was Follow-Up Long Enough for Outcomes to Occur

An acceptable length of time should be decided before quality assessment begins.

#### 4) Adequacy of Follow Up of Cohorts

This item assesses the follow-up of the exposed and non-exposed cohorts to ensure that losses are not related to either the exposure or the outcome.

Allocation of points as per rating sheet

# **Appendix G. Thresholds for Quality Assessment**

#### *Cochrane Risk of Bias (RoB) Tool* for randomized controlled trials (RCTs)

Cochrane Collaboration uses strict criteria for the threshold: A good-quality study must meet all criteria (Low RoB). A fair-quality study does not meet, or it is not clear that it meets, at least one criterion, but it has no known important limitation that could invalidate its results (Moderate RoB). A poor-quality study has important limitations and/or at least one criterion not met (High RoB).

- If all criteria met (i.e., all elements are rated as "low" risk of bias) = Good Quality
- If 1 criteria not met or 2 criteria unclear, and the assessment that this was *unlikely* to have biased the outcome, and there is no known important limitation that could invalidate the results = Moderate RoB = **Fair Quality**
- If 1 criteria not met or 2 criteria unclear, and the assessment that this was *likely* to have biased the outcome, and there are important limitations that could invalidate the results = High RoB = **Poor Quality**
- If 2 or more criteria not met = High RoB = **Poor Quality**

**Note:** Using the Cochrane RoB Tool, it is possible for a criterion to be met, even when this element was technically not part of the method; for instance, a judgment that knowledge of the allocated interventions was adequately prevented, even though there was no blinding, but the review authors judge that the outcome and the outcome measurement are not likely to be influenced by lack of blinding.

# <u>Newcastle-Ottawa Scale (N-O-S)</u> for observational studies (e.g., cohort studies and case control studies)

The Newcastle-Ottawa Scale includes 3 categories, with a maximum of 9 points, based on:

#### Selection (maximum of 4 points)

- 1) Representativeness of the exposed cohort (one point)
- 2) Selection of the non exposed cohort (one point)
- 3) Ascertainment of exposure (one point)
- 4) Demonstration that outcome of interest was not present at start of study (one point)

#### Comparability (maximum of 2 points)

- 1) Comparability of cohorts on the basis of the design or analysis
  - a) Study controls for age (one point)
  - b) Study controls for any additional factor (one point)

#### Outcome (maximum of 3 points)

- 1) Assessment of outcome (one point)
- 2) Was follow-up long enough for outcomes to occur (one point)
- 3) Adequacy of follow up of cohorts (one point)

# Scoring algorithm\*

Quality rating	# Points in Selection Domain	# Points in Comparability Domain	# Points in Outcome Domain		
Good	≥3	≥2	≥2		
Fair	2	≥1	<u>≥2</u>		
Poor	0-1	0	0-1		

# **Appendix H. Quality of Individual Studies**

Table H-1. Quality assessment of randomized controlled trials of quality improvement interventions addressing disparities in health outcomes

Author, Year	Random Sequence Generation	Allocation Concealment	Selective Reporting	Other Bias	Blinding- Participants and Personnel	Blinding- Outcome Assessment	Incomplete Outcome Data	Quality Rating
Arean et al., 2005 <sup>1</sup>	Low	Low	Low	Low	Unclear	High	Unclear	Poor
Arean et al., 2007 <sup>2</sup>	Low	Low	Low	Low	Unclear	Low	Low	Fair
Bao et al., 2011 <sup>3</sup>	Low	Low	Low	Unclear	Low	High	Unclear	Fair
Barr et al., 2001 <sup>4</sup>	Unclear	Unclear	Low	Low	High	Unclear	Unclear	Poor
Beach et al., 2007 <sup>5</sup>	Low	Low	Low	High	Low	Low	Low	Fair
Bosworth et al., 2011 <sup>6</sup>	Low	Unclear	Low	Low	Low	Low	Low	Fair
Connett and Stamler,1984 <sup>7</sup>	Low	Low	Low	Low	Low	Unclear	Low	Fair
Ferreira et. al., 2005 <sup>8</sup>	Unclear	Unclear	Low	Low	High	High	Low	Poor
Lasser et al., 20119	Low	Low	Low	Low	High	Low	Low	Fair
Miranda et al., 2003 <sup>10</sup>	Low	Low	Low	Unclear	Low	High	Low	Fair
Miranda et al., 2004 <sup>11</sup>	Low	Low	Low	Unclear	Low	High	Low	Fair
Sequist et al., 2001 <sup>12</sup>	Low	Low	Low	Low	Low	Low	Low	Good
Sherbourne et al., 2004 <sup>13</sup>	Low	Low	Low	Unclear	Low	High	Low	Fair
Siddiqui et al., 2011 <sup>14</sup>	Unclear	Unclear	Low	Low	Low	Low	Unclear	Poor
Wells et al., 2004 <sup>15</sup>	Low	Low	Low	Unclear	Low	High	High	Poor
Wells et al., 2007 <sup>16</sup>	Low	Low	High	Unclear	Low	High	High	Poor

Table H-2. Quality assessment of observational studies of quality improvement interventions addressing disparities in health outcomes

Author, Year		Representativeness of exposed cohort	Selection of the nonexposed cohort	Ascertainment of exposure	Demonstration that outcome of interest was not present at start of study	Comparability of cohorts / controlling for confounders	Assessment of outcome	Appropriate duration of followup	Adequacy of followup of cohorts	Total Score	Quality rating
	Coberley et al., 2007 <sup>17</sup>	1	1	1	1	1	1	1	0	7	Good
	Mahotiere et al., 2006 <sup>18</sup>	1	1	1	1	1	1	1	1	8	Good
	Olomu et al., 2010 <sup>19</sup>	1	1	1	0	1	1	1	0	6	Fair

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## **Appendix I. Evidence Table**

Study		Baseline		
Description Population	Intervention(s)	Characteristics	Outcomes	Disparity
Author: Condition:	Quality	Clinical:	Clinical:	Disparity before
Bao et al., 2011 Depression	improvement	NR	AD with adequate	intervention:
Region/State: Inclusion	intervention(s):	Process:	dose at 24	NR
New York City, criteria:	Collaborative care	Adequate	months*, mean	Disparity after
Philadelphia, and • MMSE score	model, including	depressant	intervention effects	intervention:
Pittsburgh ≥18	practice-based care	medication	in percentage	Differences in the
Setting: • CESD score >	managers to	dose, n (%):	points (bootstrap	intervention effects
Primary care 20 or from a 5		Collaborative	95% CI):	between the two
clinics % random	based treatment	care, any	Any college: -2.5	education groups
Enrollment sample of	recommendations,	college: 22 (34)	(-16.0 to 13.1)	did not achieve
period: those with	monitor patient	Collaborative	No college: 15.3 (2.9	statistical
May 1999- score < 20 but	clinical status, and	care, no college:		significance at
August 2001 responding	provide	46 (37)	White: 13.8 (1.8 to	months 18 and 24.
Funding: positively to	psychotherapy.	Collaborative	28.0)	Intervention
National Institute supplemental	Intervention	care, White: 49	Minority: -5.5 (-22.5	effects, difference
of Mental Health; questions	target:	(35)	to 12.2)	in mean
first author (Bao) regarding	Patients and	Collaborative	HDRS score at 24	percentage points
supported by the previous	providers	care, minority: 19	months*, mean	(bootstrap 95% CI)
Pfizer Scholar's episodes or	Groups:	(37)	intervention effects	
Grant in Health treatment of	G1: Collaborative	Usual care, any	in percentage	antidepressant
Policy. depression	care	college: 21 (32)	points (bootstrap	dose at month 24,
Conflict of Exclusion	G2: Usual care	Usual care, no	95% CI):	adjusted for
Interest: criteria:	N at enrollment:	college: 30 (33)	Any college: 1.2 (-	adequate dose at
Authors report See inclusion	NR	Usual care,	1.1 to 3.6)	baseline:
that that study <b>Disparity</b> :	N at followup:	White: 38 (39)	No college: -2.6 (-4.6	
sponsor(s) had Education level	<b>G1</b> : 214	Usual care,	to -0.4)	college, adjusted for
no role in the (No college);	<b>G2:</b> 182	minority: 13 (22)	White: -2.3 (-4.0 to -	AD with adequate
design and Race/ ethnicity	Any college:	Collaborative	0.1)	dose at baseline:
conduct of the (Minority)	<b>G1</b> : 73	care, no college:	Minority: 1.2 (-1.4 to	
study. Referent group	: <b>G2</b> : 73	$21.2 \pm 5.3$	4.2)	Minority – White,
Internal; Any	No college:	Collaborative		adjusted for AD with
<b>Design:</b> college; White	<b>G1</b> : 141	care, White: 21.0		adequate dose at
RCT (secondary Subgroup or	<b>G2:</b> 182	± 5.7		baseline: -19.3 (-
analysis) secondary	White:	Collaborative		40.6 to 3.3)
analysis	<b>G1:</b> 151	care, minority:		Intervention
description:	<b>G2</b> : 111	$20.5 \pm 5.6$		effects, difference
Randomization	Minority:	Usual care, any		in mean
not stratified by	<b>G1</b> : 63	college: 19.1 ±		percentage points
education level	<b>G2</b> : 71	5.3		(bootstrap 95% CI)
or race/ethnicity	Length of	Usual care, no		for HDRS score at
•	followup:	college: 20.0 ±		month 24:
	24 months	5.6		No college – some
	Measure of	Usual care,		college: -3.8 (-6.8 to
	fidelity:	White: 19.5 ± 5.7		-0.4)
	NR	Usual care,		Minority – White:
		minority: 19.6 ±		3.5
*Intervention offects based on mixed	offeets legistic model vi	5.1		(-0.1 to 6.9)

<sup>\*</sup>Intervention effects based on mixed effects logistic model with random intercepts at the patient level

Study		diffico in ficaltif o	Baseline	,	_
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
	•			Clinical:	Disparity before
Author: Bosworth et al.,	Condition: Hypertension	Quality improvement	Clinical: Systolic blood	Systolic blood	intervention:
2011	Inclusion	interventions:		•	NR
Region/State:	criteria:	Tailored behavioral	pressure, mean mm Hg ± SD:	pressure, 12 or 24 months:	Disparity after
Durham, North		self-management	G1a+G2a+G3a+	G1a: NR	intervention:
Carolina	<ul> <li>Hypertension diagnosis for &gt;</li> </ul>	(G1)	G4a:	G1b: NR	NR
Setting:	12 months	Home blood	121.5 ± 15.6	G2a: NR	Change in
University-		pressure monitoring			disparity:
affiliated general		(G2)	G4b:	G3a: NR	Differences in
internal	philliary care	Combination of the	128.3 ± 19.0	G3b: NR	systolic blood
medicine clinics	study clinic	two interventions	Diastolic blood	G4a: NR	pressure, mm Hg
Enrollment	Received	(G3)	pressure, mean	G4b: NR	(95% CI), 12
period:	hypertensive	Usual care (G4)	mm Hg ± SD:		months:
May 2001-	medication	Intervention	G1a+G2a+G3a+	Diastolic blood	G1a vs. G4a:
December 2002	prescription in	target:	G4a:	pressure, 12 or 24	2.3 (-2.4 to 7.0)
Funding:	the previous	Patient blood	68.8 ± 10.6	months:	G2a vs. G4a:
Department of	year	pressure changes	G1b+G2b+G3b+		-1.5 (-6.1 to 3.2)
Veterans Affairs	<ul> <li>Scheduled for a</li> </ul>	·_	G4b:	G1b: NR	G3a vs. G4a:
Health Services	primary care	G1a: White,	73.7 ± 10.5	G2a: NR	-0.7 (-5.2 to 3.9)
Research	physician	behavioral		G2b: NR	G1b vs. G4b:
Division;	appointment	intervention		G3a: NR	-5.7 (-10.0 to -1.4)
National	within next 30	G1b: nonwhite,		G3b: NR	G2b vs. G4b:
Institutes of	days	behavioral		G4a: NR	-5.5 (-10.3 to -0.8)
Health	Residing in one	intervention		G4b: NR	G3b vs. G4b:
Conflict of	of 32 specified	G2a: White, home			-5.3 (-10.1 to -0.5)
Interest:	zip codes	blood pressure		Race x time x	
NR	Exclusion	monitor intervention		treatment group	Differences in
Design:	criteria:	G2b: nonwhite,		effect suggested	systolic blood
RCT	<ul> <li>Diagnosis of</li> </ul>	home blood		likely differential	pressure, mm Hg
	dementia,	pressure monitor		intervention effects	(95% CI), 24
	Parkinson's	intervention		over time for White	months:
	disease, atrial	G3a: White,		and nonwhite	G1a vs. G4a:
	fibrillation, or	combined		patients for both	2.5 (-3.1 to 8.1)
	end-stage renal	intervention		systolic blood	G2a vs. G4a:
	disease	<b>G3b</b> : nonwhite,		pressure (p=0.08)	-0.8 (-6.4 to 4.7)
	<ul> <li>Residing in a</li> </ul>	combined		and diastolic blood	G3a vs. G4a:
	nursing home	intervention		pressure (p=0.01).	-0.4 (-5.9 to 5.0) <b>G1b vs. G4b:</b>
	or receiving	<b>G4a</b> : White, usual			
	home health	care			-0.6 (-6.0 to 4.8) <b>G2b vs. G4b:</b>
	care	<b>G4b</b> : nonwhite, usual care			
		N at baseline:			1.3 (-4.4 to 7.1) <b>G3b vs. G4b</b> :
		<b>G1a</b> : 69			-7.5 (-13.7 to -1.4)
		<b>G1b</b> : 89			-1.0 (-10.1 lU -1.4)
		<b>G2a</b> : 78			
		<b>G2b:</b> 78			
		<b>G3a</b> : 88			
		<b>G3b</b> : 70			
		<b>G4a</b> : 70			
		<b>G4b:</b> 87			
		<b></b> .			

	addressing disp	anties in neath c	Baseline	eu)	
Study Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Bosworth et al.,	Exclusion	N at 12 months	Characteristics	Outcomes	Differences in
2011	criteria (cont):	followup:			diastolic blood
(continued)	Hospitaliza-tion	•			pressure, mm Hg
(continuou)	for stroke or	<b>G1b</b> : 71			(95% CI), 12
	heart attack,	<b>G2a:</b> 68			months:
	surgery for	<b>G2b:</b> 50			G1a vs. G4a:
	blocked	<b>G3a:</b> 75			2.9 (0.4 to 5.4)
	arteries, or	<b>G3b:</b> 47			G2a vs. G4a:
	diagnosed w/	<b>G4a</b> : 60			0.1 (-2.4 to 2.6)
	metastatic	<b>G4b</b> : 71			G3a vs. G4a:
	cancer in	N at 24 months			1.3 (-1.2 to 3.7)
	previous 3 mo	followup:			G1b vs. G4b:
	Poor vision or	<b>G1a:</b> 62			-3.3 (-5.6 to -0.9)
	difficulty	<b>G1b</b> : 62			G2b vs. G4b:
	hearing on the	<b>G2a</b> : 64			-3.7 (-6.2 to -1.1)
	telephone	<b>G2b</b> : 49			G3b vs. G4b:
	<ul> <li>Difficulty</li> </ul>	<b>G3a:</b> 72			-2.7 (-5.3 to -0.2)
	understanding	<b>G3b:</b> 38			
	English	<b>G4a</b> : 60			Differences in
	<ul> <li>Participation in</li> </ul>	<b>G4b</b> : 68			diastolic blood
	another BP	Length of			pressure, mm Hg
	study	followup:			(95% CI), 24
	<ul> <li>Spouse in the</li> </ul>	24 months			months:
	current study	Measure of			G1a vs. G4a:
	<ul><li>Arm</li></ul>	fidelity:			2.0 (-1.2 to 5.1)
	circumference	NR			G2a vs. G4a:
	>17 in or wrist				0.1 (-3.0 to 3.2)
	circumference				G3a vs. G4a:
	>8.5 in				0.5 (-2.5 to 3.5)
	Disparity				G1b vs. G4b:
	indicator(s):				0.6 (-2.4 to 3.6) <b>G2b vs. G4b:</b>
	Race/ethnicity				-0.6 (-3.9 to 2.6)
	(Nonwhite)				G3b vs. G4b:
	Referent group:				-3.5 (-7.0 to -0.1)
	Internal; White				-3.3 (-7.0 to -0.1)
	Subgroup or				
	secondary				
	analysis				
	description:				
	Post hoc analysis				
	of an RCT <b>White, n:</b>				
	308				
	Nonwhite*, n:				
	328				
	J20				

Baseline mean systolic blood pressure reported for White and nonwhite groups.

<sup>\*</sup>Majority of the nonwhite group were African American (95%)

The overall race by time by treatment group effect (6df test) suggested likely differential intervention effects over time for White and nonwhite patients for both systolic blood pressure (p=08) and diastolic blood pressure (p=.01).

Study	addressing dispa	THOO III HOURTH	Baseline	iladaj	
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Author:	Condition:	Quality	Clinical:	Clinical:	Disparity before
Lasser et al.,	Colorectal cancer	improvement	NA	NA	intervention:
2011	(CRC) screening	intervention:	Process:	Process:	NA; eligible
Region/State:	Inclusion criteria:	•		Colorectal cancer	participants did not
Massachusetts	<ul> <li>Patients aged 52</li> </ul>	based	screening:	screen test	have CRC
Setting:	to 74 years	intervention	Eligible	completed, n (%):	screening
Primary care	<ul> <li>Spoke English,</li> </ul>	Intervention	participants did	<b>G1a</b> : 38 (33.9)	Disparity after
practice-based research	Haitian Creole,	target: Patients	not have a CRC screening test	<b>G1b</b> : 25 (39.7) <b>G1c</b> : 30 (26.8)	intervention: Intervention was
network	Portuguese, or Spanish as	Groups:	screening test	<b>G1d</b> : 49 (39.8)	particularly
Enrollment	primary language	G1: participants		<b>G2a:</b> 18 (16.5)	beneficial for non-
period:	Not completed	randomized to		<b>G2b</b> : 11 (16.7)	English language
September	CRC screening*	intervention		<b>G2c</b> : 24 (21.4)	participants
2008-March	Exclusion criteria:	<b>-</b> 4 1.11		<b>G2d</b> : 22 (18.6)	
2009	Patients with	participants,		Colorectal cancer	
Funding:	acute illness, end	intervention		screening test	
American	stage renal	G1b: black		completed,	
Cancer Society	disease, severe	participants,		intervention vs.	
Conflict of	psychiatric	intervention		control, %, p-value:	
Interest: None	condition, active	G1c: English		<b>G1a vs. G2a</b> : 17.4,	
Design:	substance abuse,	speaking participants,		p=0.003 <b>G1b vs. G2b</b> : 23.0,	
RCT	or cognitive	intervention		p=0.004	
NO I	impairment <b>Disparity</b>	G1d: non-English		G1c vs. G2c: 5.4,	
	indicator(s):	speaking		p=0.35	
	Race	participants,		G1d vs. G2d: 21.2,	
	Primary language	intervention		p<0.001	
	Referent group:	G2: participants			
	Internal	randomized to			
	Subgroup or	usual care			
	secondary	G2a: white			
	analysis	participants,			
	description:	usual care			
	Stratified analysis	<b>G2b:</b> black participants,			
	according to	usual care			
	primary language,	G2c: English			
	age, race, and health insurance	speaking			
	status.	participants,			
	otatuo.	usual care			
		G2d: non-English			
		speaking			
		participants,			
		usual care			

Study			Baseline	•	
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Lasser et al.,		N at enrollment:			
2011		<b>G1a</b> : 112			
(continued)		<b>G1b</b> : 63			
		<b>G1c</b> : 112			
		<b>G1d</b> : 123			
		<b>G2a:</b> 109			
		<b>G2b:</b> 66			
		<b>G2c</b> : 112			
		<b>G2d</b> : 118			
		N at follow-up:			
		<b>G1a</b> : 112			
		<b>G1b</b> : 63			
		<b>G1c</b> : 112			
		<b>G1d</b> : 123			
		<b>G2a:</b> 109			
		<b>G2b</b> : 66			
		<b>G2c</b> : 112			
		<b>G2d:</b> 118			
		Length of			
		follow-up:			
		Up to one year			
		after enrollment			
		Measure of			
		fidelity:			
		<ul> <li>Navigators</li> </ul>			
		contacted 181 of			
		the 235			
		intervention			
		patients (77%).			

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

	addressing dispa	inies in neann ou	Baseline	ueu)	
Study Description	Population	Intervention(s)	Characteristics	Outcomes	Dieparity
Author:	Condition:		Clinical:	Clinical:	Disparity Disparity before
	Colorectal cancer	Quality improvement	NA	NA	intervention:
Siddiqui et al., 2011	(CRC) screening	intervention:			
Region/State:	Inclusion criteria:		Process:	Process:	NA; eligible
Pennsylvania	Male and female	and reminders	Eligible participants did	CRC screening test completed during	have CRC
Setting:	patients aged 50-		not have CRC	the 12 month	screening
Academic	74 years	messages and	screening	observation period:	
primary care	<ul> <li>Had one visit to</li> </ul>	phone reminders	Screening	<b>G1a</b> : 32.9	intervention:
practice	the Jefferson	from health		<b>G1b</b> : 32.1	Screening rate for
Enrollment	Family Medicine	educators		<b>G2a</b> : 55.1	whites compared to
period:	Associates	Intervention		<b>G2b</b> : 40.78	blacks, OR (95%
February 2002-	practice with the	target:		<b>G3a</b> : 50.4	CI) p value:
March 2002	previous two	Appropriate		<b>G3b</b> : 40.2	<b>G1a vs. G1b</b> : 1.01
Funding:	years	screening for		<b>G4a</b> : 53.9	(0.64 to 1.61)
	Had complete	colorectal cancer		<b>G4b</b> : 47.7	p=0.956
Institute and	contact	Groups:			<b>G2a vs. G2b</b> : 1.68
Pennsylvania	information	G1: participants			(1.10 to 2.58)
Department of	available	randomized to			p=0.017
Health	<ul> <li>Exclusion</li> </ul>	control			<b>G3a vs. G3b</b> : 1.42
Conflict of	criteria:	G1a: white, control			(0.92 to 2.21)
Interest:	<ul> <li>No prior</li> </ul>	<b>G1b</b> : black, control			p=0.117
None	diagnosis of	<b>G2:</b> participants			<b>G4a vs. G4b</b> : 1.25
Design:	colorectal	randomized to			(0.81 to 1.92)
RCT	neoplasia or	standard			p=0.316
	inflammatory	intervention			G2a + G3a vs. G2b
	bowel disease	<b>G2a:</b> white, standard			<b>+ G3b</b> : 1.56 (1.14 to 2.12) p=0.005
	<ul> <li>Had not</li> </ul>	into mantion			G2a + G3a +G4a
	undergone recent	G2b: black,			vs. G2b + G3b +
	CRC screening*	standard			<b>G4b</b> : 1.44 (1.12 to
	Disparity	intervention			1.86) p=0.005
	indicator(s): Race	G3: participants			, р
	Referent group:	randomized to			
	Internal	tailored intervention			
	Subgroup or	G3a: white, tailored			
	secondary	intervention			
	analysis	G3b: black, tailored			
	description:	intervention			
	Post-hoc analysis	<b>G4</b> : participants			
	of race-intervention	randomized to			
	effects from a RCT	tailored intervention			
	of tailored	plus reminder			
	interventions for	phone call			
	CRC screening	<b>G4a:</b> white, tailored			
		intervention plus reminder call			
		<b>G4b:</b> black, tailored			
		intervention plus			
		reminder call			
		TOTTINION CAN			

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Siddiqui et al.,		N at enrollment:			
2011		<b>G1a</b> : N/R			
(continued)		<b>G1b</b> : N/R			
,		<b>G2a</b> : N/R			
		<b>G2b</b> : N/R			
		<b>G3a</b> : N/R			
		<b>G3b</b> : N/R			
		<b>G4a</b> : N/R			
		<b>G4b</b> : N/R			
		N at follow-up:			
		<b>G1a</b> : 146			
		<b>G1b</b> : 215			
		<b>G2a</b> : 156			
		<b>G2b</b> : 206			
		<b>G3a</b> : 135			
		<b>G3b</b> : 214			
		<b>G4a</b> : 141			
		<b>G4b</b> : 217			
		Length of follow-			
		up:			
		12 months			
		Measure of			
		fidelity:			
		N/R			

	interventions addressing disparities in health outcomes (continued)						
Study			Baseline				
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity		
Author:	Condition:	Quality	Clinical	Clinical (inpatient	Disparity before		
Olomu et al.,	Acute myocardial	-	(inpatient	events), n (%):	intervention:		
2010	infarction (AMI)	intervention(s):	events), n (%):	Hypotension:	NR		
Region/State:	Inclusion	American College	Hypotension:	<b>G2a</b> : 410 (33.9)	Disparity after		
Michigan	criteria:	of Cardiology's	<b>G1a:</b> 350 (30.2)	<b>G2b</b> : 87 (31.1)	intervention:		
Setting:	• Pre-GAP	Guidelines Applied	<b>G1b</b> : 68 (32.4)	Shock:	Clinical (inpatient		
Academic and	sample: 50%	in Practice (GAP)	Shock:	<b>G2a</b> : 16 (1.3)	events), OR (95%		
community	random sample	program including	<b>G1a:</b> 14 (1.2)	<b>G2b</b> : 1 (0.7)	CI)*		
hospitals	with ≥ 20 cases		<b>G1b</b> : 2 (1.0)	Heart failure or	Hypotension:		
Enrollment	per hospital of	orders; pocket	Heart failure or	pulmonary edema:	<b>G2b vs. G2a:</b> 0.88		
period:	Medicare AMI	guideline;	pulmonary	<b>G2a:</b> 529 (43.8)	(0.66 to 1.16)		
NR Fundings	patients	standardized	edema:	<b>G2b:</b> 137 (48.9)	Shock:		
Funding: National	(principal	discharge tool;	<b>G1a:</b> 548 (47.3) <b>G1b:</b> 104 (49.5)	Stroke: <b>G2a:</b> 56 (4.6)	<b>G2b vs. G2a</b> : 0.54		
American	diagnosis code		Stroke:	<b>G2b:</b> 20 (7.1)	(0.12 to 2.35) Heart failure or		
College of	410.xx) treated at participating	patients, physicians		Renal insufficiency:	pulmonary edema:		
Cardiology	hospitals in the	and nurses; system		<b>G2a:</b> 259 (21.6)	<b>G2b vs. G2a:</b> 1.23		
Foundation;	year before	for measurement	Renal	<b>G2b</b> : 97 (35.3)	(0.95 to 1.60)		
Centers for	GAP	Intervention	insufficiency:	Hemorrhage/	Stroke:		
Medicare and	implementa-	target:	<b>G1a:</b> 266 (23.1)	bleeding	<b>G2b vs. G2a:</b> 1.58		
Medicaid	tion	provision of	<b>G1b:</b> 73 (34.9)	<b>G2a:</b> 291 (24.1)	(0.93 to 2.69)		
Services;	Post-GAP	evidence-based	Hemorrhage/	<b>G2b:</b> 85 (30.3)	Renal insufficiency:		
Michigan Peer	sample: 95-	treatments in	bleeding	Process, in-	<b>G2b vs. G2a:</b> 1.98		
Review	100% sampling	hospital (aspirin,	<b>G1a</b> : 271 (23.4)	hospital	(1.49 to 2.63)		
Organization;	of all Medicare	beta-blocker) and	<b>G1b:</b> 85 (30.3)	procedures/	Hemorrhage/		
Pfizer;	AMI patients in	at discharge	Process, in-	treatment, n (%):	bleeding:		
AstraZeneca;	the 4 months	(aspirin, beta-	hospital	Echocardiogram:	<b>G2b vs. G2a:</b> 1.38		
<b>Greater Detroit</b>	immediately	blocker, lipid-	procedures/	<b>G2a:</b> 716 (59.1)	(1.03 to 1.83)		
Area Health	after GAP	lowering agents,	treatment, n	<b>G2b:</b> 171 (61.1)	Process, in-hospital		
Council;	implementa-	angiotensin-	(%):	Cardiac	procedures/		
Mardigian	tion at each	converting enzyme	Admission tool	catheterization:	treatment, OR (95%		
Foundation;	hospital	inhibitor, and	used:	<b>G2a:</b> 614 (50.8)	CI)*		
University of	Exclusion	smoking cessation	<b>G1a:</b> 102 (8.8)	<b>G2b:</b> 103 (36.8)	Echocardiogram:		
Michigan Ann	criteria:	counseling), in-	<b>G1b:</b> 13 (6.2)	PCI:	<b>G2b vs. G2a:</b> 1.08		
Arbor	See inclusion	hospital	Echocardio-	<b>G2a:</b> 309 (25.6)	(0.83 to 1.41)		
Conflict of	criteria	complications	gram:	<b>G2b:</b> 37 (13.2)	Cardiac		
Interest:	Disparity:		<b>G1a:</b> 715 (61.7)	CABG:	catheterization:		
NR Da a la succession	Race/ethnicity		<b>G1b:</b> 128 (61.0)	<b>G2a:</b> 127 (10.5)	<b>G2b vs. G2a:</b> 0.56		
Design:	(Nonwhite)		Cardiac	<b>G2b</b> : 14 (5.0)	(0.43 to 0.74)		
Retrospective	Referent group:		catheterization:		PCI:		
cohort	Internal; White		<b>G1a:</b> 527 (45.5)		<b>G2b vs. G2a:</b> 0.44		
			<b>G1b:</b> 94 (44.8) PCI:		(0.31 to 0.64) CABG:		
			<b>G1a:</b> 229 (19.8)		<b>G2b vs. G2a:</b> 0.46		
			<b>G1b:</b> 33 (15.7) CABG:		(0.26 to 0.79)		
			<b>G1a:</b> 101 (8.7)		Aspirin administered:		
			<b>G1b:</b> 10 (4.8)		<b>G2b vs. G2a:</b> 0.73		
			<b>GID.</b> 10 (4.0)		(0.37 to 1.47)		
					(0.07 to 1.47)		

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Description Olomu et al., 2010 (continued)	Subgroup or secondary analysis description: Comparison of nonwhites vs. Whites during the post-guideline implementation period Measure of fidelity: Admission tool used: G2a: 551 (45.6) G2b: 120 (42.9) OR 0.90 (0.69 to 1.16) Discharge tool used: G2a: 364 (30.1) G2b: 66 (23.6)	Groups: G1a: White, patients before GAP G1b: nonwhite, patients before	Aspirin administered: G1a: 425 (85.7) G1b: 73 (81.1) Beta-blocker initiated: G1a: 192 (76.8) G1b: 35 (66.0) Process, discharge procedures/ treatment, n (%): Discharge tool used: G1a: 21 (1.8) G1b: 2 (1.0) Aspirin prescribed: G1a: 296 (83.4) G1b: 58 (89.2) Beta-blocker prescribed: G1a: 102 (87.9) G1b: 31 (96.9) Lipid-lowering therapy prescribed: G1a: 163 (80.3) G1b: 24 (77.4) ACE-inhibitor prescribed: G1a: 164 (82.8) G1b: 29 (93.6) Counseling for smoking cessation: G1a: 48 (39.4)	Aspirin administered: G2a: 419 (92.9) G2b: 115 (90.6)Beta-blocker initiated: G2a: 192 (82.4) G2b: 45 (83.3) Process, discharge procedures/ treatment, n (%): Aspirin prescribed: G2a: 393 (94.2) G2b: 74 (90.2) Beta-blocker prescribed: G2a: 129 (94.9) G2b: 20 (90.9) Lipid-lowering therapy prescribed: G2a: 230 (83.0) G2b: 45 (79.0) ACE-inhibitor prescribed: G2a: 159 (86.9) G2b: 31 (93.9) OR 2.34 (0.53 to 10.41) Counseling for smoking cessation: G2a: 116 (73.4) G2b: 24 (50.0) Admission tool used: G2a: 364 (30.1) G2b: 66 (23.6)	Beta-blocker initiated:  G2b vs. G2a: 1.07 (0.48 to 2.36) Process, discharge procedures/ treatment, OR (95% CI)* Aspirin prescribed: G2b vs. G2a: 0.56 (0.24 to 1.31) Beta-blocker prescribed: G2b vs. G2a: 0.54 (0.11 to 2.80) Lipid-lowering therapy prescribed: G2b vs. G2a: 0.77 (0.38 to 1.56) ACE-inhibitor prescribed: G2b vs. G2a: 2.34 (0.53 to 10.41) Counseling for smoking cessation: G2b vs. G2a: 0.36 (0.19 to 0.71) Admission tool used: G2b vs. G2a: 0.90 (0.69 to 1.16) Discharge tool used: G2b vs. G2a: 0.72

<sup>\*</sup>Includes post-GAP patients only, nonwhite vs. White; unclear how the authors derived the OR

Comments:

The intent of the intervention was not to reduce disparity. The intention of the intervention was to improve overall care for patients. The odds ratios focus on whether there was a post-GAP implementation – no demonstration of a pre-GAP disparity in the data set and no analysis of whether a disparity was closed via the intervention.

<sup>†</sup> N at followup is equal to N at enrollment because this is retrospective cohort study.

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study			Baseline	• .	
Description	Population	Intervention(s)	Characteristics		Disparity
Study Description  Author: Sequist et al., 2010 Region/State: Massachusetts Setting: Community practice; Harvard Vanguard Medical Associates Enrollment period: June 2007-May 2008 Funding: Robert Wood Johnson Foundation Conflict of Interest: First author discloses consultancy for Aetna Design: Cluster RCT	Population  Condition: Diabetes mellitus Inclusion criteria: • Chronic disease focused primary care teams comprising physicians, nurse practitioners (NPs) and/or physician assistants (PAs) from 8 health centers • Primary care teams were caring for patients with diabetes, defined as having problem list diagnosis of diabetes in the electronic medical record and at least one of the following: fasting plasma glucose >3.3 mmol/L; random plasma glucose >5.2 mmol/L; completed HbA1c test. Exclusion criteria: NP	intervention(s): Cultural competency training of primary care teams over two days for NPs and PAs and one day for physicians addressed racial and cultural biases, appropriate methods of collecting clinically relevant cultural data, and ways to incorporate cultural data into patient care plans. Training consisted of lectures, group discussion and community engagement activity through which clinicians met with Black patients with diabetes and learned about barriers to diabetes management. Monthly race- stratified physician level diabetes performance feedback reports were provided. Intervention target: Providers; outcomes	Clinical: HbA1c <7 percent, n (%): G1a+G2a: 1080 (40) G1b+G2b: 2241 (46.1) p<0.001 LDL cholesterol <2.59 mmol/L, n (%): G1a+G2a: 1170 (43.4) G1b+G2b: 2686 (55.3) p<0.001 Blood pressure <130/80 mm Hg, n (%): G1a+G2a: 640 (23.7) G1b+G2b: 1535 (31.5) p<0.001 Process: Annual HbA1c exam, n (%): G1a+G2a:2395 (88.7) G1b+G2b:4230 (87.1) p=0.139 Annual LDL cholesterol exam, n (%): G1a+G2a:2246 (83.2) G1b+G2b:4032 (83.0) p=0.99 Annual blood pressure	LDL cholesterol <2.59 mmol/L, %: G1a: 49.0 G1b: 61.6 G2a: 50.8 G2b: 59.6 Blood pressure <130/80 mm Hg, %: G1a: 24.0 G1b: 30.5 G2a: 25.4	p<0.001 LDL cholesterol <2.59 mmol/L: p<0.001 Blood pressure <130/80 mm Hg: p<0.001 Differences in process measures of disease for White and Black patients, p value: Annual HbA1c exam: p=0.139 Annual LDL cholesterol exam: p=0.99 Annual blood pressure measurement: p=0.035 Disparity after intervention Intervention Intervention effects on difference between White and Black patients*, p value: HbA1c <7 percent: p=0.22 LDL cholesterol <2.59 mmol/L:
	NR Disparity: Race/ethnicity (Black) Referent group: Internal; White	measured for patients	measurement, n (%): G1a+G2a:2561 (94.9) G1b+G2b:4575 (94.2) p=0.035		p=0.34 Blood pressure <130/80 mm Hg: p=0.68

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Sequist et al., 2010 (continued)	Subgroup or secondary analysis description: NA	Groups: G1a: Black patients, intervention G1b: White patients, intervention G1c: primary care team, intervention G2a: Black patients, control G2b: White patients, control G2c: primary care team, control N at enrollment: G1a: 1401 G1b: 2383 G2a: 1298 G2b: 2475 G1c:15 teams; 46 physicians; 16 NPs or PAs G2c: 16 teams; 45 physicians; 17 NPs or PAs N at followup: G1a: 1401 G1b: 2383 G2a: 1298 G2b: 2475 Length of followup: 12 months Measure of fidelity: NA			

<sup>\*</sup>After adjustment for clustering by primary care team.

The primary care teams were chronic disease management-focused primary care teams and it was 2-3 primary care physicians working collaboratively with a NP or PA; the NP or PA was primary responsible for diabetes management.

The monthly feedback reports highlighted Black-White differences in rates of achieving ideal control of HbA1c, LDL and BP within each clinician's patient panel and across the 8 health centers. Teams also received detailed, patient-specific report during months 4 and 9.

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

	addressing disp	danties in neatth o	·		
Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
			Clinical:		
Author:	Condition: Depression	Quality improvement		Clinical outcomes at 12 months:	Disparity before intervention:
Arean et al., 2007	Inclusion	intervention(s):	SCL-20 score, mean ± SD:	SCL-20 score,	General health,
		Collaborative care		mean ± SD:	mean ± SD:
Region/State: 5 U.S. states	•	model, including	G1a+G2a:	<b>G1a:</b> 1.07 ± 0.71	G1a+G2a:
	<ul> <li>60 years or older</li> </ul>	stepped care	$1.7 \pm 0.6$		
Setting: Primary care		approach to	G1b+G2b:	<b>G1b:</b> 0.95 ± 0.65	$3.6 \pm 1.0$
clinics	<ul> <li>Met criteria for major</li> </ul>	managing depression; primary	$1.7 \pm 0.6$	<b>G2a:</b> 1.45 ± 0.66	G1b+G2b:
Enrollment	depression or	care provider	p=0.59	<b>G2b:</b> $1.36 \pm 0.68$	3.2 ± 1.1
period:	dysthymia	education;		G1a vs. G2a,	p<0.01
NR	according to	depression clinical	General health,	adjusted OR (95%	
Funding:	SCID	specialist/	mean ± SD:	CI, p-value): -0.39 (-0.50, -0.27,	PCS-12, mean ±
John A. Hartford	Exclusion	depression care manager works with	G1a+G2a:	p<0.001)	SD:
Foundation;	ontona.	patient and primary	$3.6 \pm 1.0$	G1b vs. G2b,	G1a+G2a:
California	See inclusion	care provider,	G1b+G2b:	adjusted OR (95%	$38.6 \pm 7.1$
Healthcare	Disparity:	medication	3.2 ± 1.1	CI, p-value): -0.41	G1b+G2b:
Foundation;	Socioeconomic	monitoring and brief		(-0.49, -0.33,	41.1 ± 7.4
Hogg Foundation;	status (Poor, <	psychotherapy; use of clinical	p<0.01	p<0.001)	p<0.01
Robert Wood	30% of area median income)	information tracking	DOO 40	General health,	Disparity after
Johnson	Referent group:	system; ready	PCS-12, mean ± SD:	mean ± SD:	intervention:
Foundation	•	access to a	G1a+G2a:	<b>G1a:</b> 3.40 ± 1.01	No interaction
Conflict of	Internal; Not poor, living >	psychiatrist		<b>G1b:</b> $3.06 \pm 0.98$	between income
Interest:	30% of area	Intervention	$38.6 \pm 7.1$	<b>G2a:</b> 3.69 ± 0.98	status and use of
NR	median income	target: Appropriate	G1b+G2b:	<b>G2b:</b> 3.38 ± 0.97	depression
Design:	Subgroup or	management of depression	41.1 ± 7.4	G1a vs. G2a,	treatment, satisfaction or other
RCT (secondary	· . ·	-	p<0.01	adjusted OR (95%	clinical outcomes;
analysis)	analysis	Groups:	Process, n (%):	CI, p-value): -0.29 (-0.45, -0.12,	no interaction
	description:	<b>G1a:</b> Collaborative	Any	(-0.45, -0.12, p=0.001)	between treatment
	Randomization not stratified by	care, poor <b>G1b:</b> Collaborative	antidepressant	G1b vs. G2b,	condition and time
	income; post hoc	care, not poor	use in last 3	adjusted OR (95%	with service use or
	subgroup	<b>G2a:</b> Usual care,	months:	CI, p-value): -0.32	other outcomes in mixed effects
	analysis	poor	G1a+G2a:	(-0.43, -0.21,	models
	conducted to	<b>G2b:</b> Usual care,	243 (42)	p<0.001)	
	compare outcomes in each		G1b+G2b:		Magnitude of
	income group	N at enrollment:	528 (43)		intervention effects
	between	<b>G1a+G2a</b> : 576	p=0.71		similar across
	intervention and	<b>G1b+G2b</b> : 1225	Any specialty		income groups.
	control	N at followup:	mental health		
		NR	visits of		
		IVIX	psychotherapy in the last 3 months:		
			G1a+G2a:		
			46 (8)		
			G1b+G2b:		
			105 (9)		
			p=0.68		

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Arean et al., 2007 (continued)		Length of followup: 12 months Measure of fidelity: NR	Satisfaction with depression care (rating excellent or very good): G1a+G2a: 104 (50) G1b+G2b: 200 (51) p=0.76	PCS-12, mean ± SD: G1a: 177 (65) G1b: 397 (65) G2a: 137 (48) G2b: 283 (49) G1a vs. G2a, adjusted OR (95% Cl, p-value): 1.46 (0.33, 2.60, p=0.011)G1b vs. G2b, adjusted OR (95% Cl, p-value): 1.67 (0.78, 2.55, p<0.001)  Satisfaction with depression care (rating excellent or very good), n (%): G1a: 168 (71) G1b: 414 (78) G2a: 89 (43) G2b: 182 (50) G1a vs. G2a, adjusted OR (95% Cl, p-value): 3.07 (1.23, 7.65, p=0.026)  G1b vs. G2b, adjusted OR (95% Cl, p-value): 3.35 (1.94, 5.77, p<0.001)	

Study	<b>-</b>		Baseline	•	<b>.</b>
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Arean et al.,				Process outcomes	
2007 continued)				at 12 months, n (%):	
continuea)				Any antidepressant:	
				<b>G1a:</b> 177 (65)	
				<b>G1b</b> : 397 (65)	
				<b>G2a:</b> 137 (48)	
				<b>G2b:</b> 283 (49)	
				G1a vs. G2a,	
				adjusted OR (95%	
				CI, p-value): 3.25	
				(2.14, 4.96,	
				p<0.001)	
				G1b vs. G2b,	
				adjusted OR (95%	
				CI, p-value): 2.17	
				(1.53, 3.08,	
				p<0.001)	
				Any psychotherapy: <b>G1a:</b> 104 (40)	
				<b>G1b:</b> 265 (44)	
				<b>G2a:</b> 42 (15)	
				<b>G2b</b> : 91 (16)	
				G1a vs. G2a,	
				adjusted OR (95%	
				CI, p-value): 4.16	
				(2.52, 6.85,	
				p<0.001)	
				C4h va C2h	
				G1b vs. G2b, adjusted OR (95%	
				CI, p-value): 4.33	
				(3.14, 5.97,	
				p<0.001)	
				p (0.001)	
				Any depression	
				treatment:	
				<b>G1a:</b> 200 (77)	
				<b>G1b</b> : 474 (78)	
				<b>G2a:</b> 152 (53)	
				<b>G2b</b> : 309 (54)	
				G1a vs. G2a,	
				adjusted OR (95%	
				CI, p-value): 3.80	
				(2.46, 5.85,	
				p<0.001)	
				G1b vs. G2b,	
				adjusted OR (95%	
				CI, p-value): 3.58	
				(2.59, 4.95,	
				p<0.001)	

Study	addressing di	sparities in near	Baseline	nucu)	
Description	Population	Intervention(		Outcomes	Disparity
Author:	Condition:	Quality	Clinical:	Clinical:	Disparity before
Beach et al.,	Cancer	improvement	NA	NA	intervention:
2007	screening	intervention(s):	Process:	Process:	Up-to-date cancer
Region/State:		Scripted	Up-to-date cancer	Up-to-date cancer	screening status*
New York City,		telephone support	screening status*	screening status§	
New York		provided by a			Breast cancer, OR
Setting:		Prevention Care		Breast cancer, n (%):	(95%):
	<ul> <li>Received care</li> </ul>	Manager (PCM)	<b>G1a+G2a</b> : 264/498	<b>G1a</b> : 144/239 (60)	G1a+G2a vs.
Migrant Health	101 = 0 111011410		(53.0)	<b>G1b</b> : 310/431 (72)	<b>G2a+G2b</b> : 1.46
Centers	at a C/MHC	target:	<b>G1b+G2b</b> : 528/848	<b>G2a</b> : 144/259 (56)	(1.16-1.84),
(C/MHC)	<ul> <li>Not up to date</li> </ul>	Patient behavior	(62.3)	<b>G2b</b> : 243/417 (58)	p<0.001
Enrollment	for breast,	Groups:	0	0	0
period:	cervical, or	G1a: English	Cervical cancer†, n	Cervical cancer†, n	Cervical cancer†,
November	colorectal	speaking,	(%): <b>G1a+G2a:</b> 209/368	(%): <b>G1a</b> : 106/181 (59)	OR (95%): <b>G1a+G2a vs.</b>
2001 - October 2002	caricci	intervention <b>G1b</b> : Spanish	(56.8)	<b>G1b</b> : 237/310 (76)	G2a+G2b: 1.19
Funding:	screening	speaking,	<b>G1b+G2b</b> : 365/599	<b>G2a</b> : 100/187 (53)	(0.90-1.56)
National	Preferred	intervention	(60.9)	<b>G2b</b> : 173/289 (60)	(0.90-1.50)
Cancer	language was	G2a: English	(00.9)	<b>G2B</b> . 173/203 (00)	Colorectal cancer‡,
Institute	English,	speaking, usual	Colorectal cancert in	Colorectal cancer‡, n	
Conflict of	Spanish or Haitian Creole		(%):	(%):	G1a+G2a vs.
Interest:	Exclusion	<b>G2b</b> : Spanish	<b>G1a+G2a</b> : 94/394	<b>G1a</b> : 95/190 (50)	<b>G2a+G2b:</b> 0.83
NR	criteria:	speaking, usual	(23.9)	<b>G1b</b> : 184/338 (54)	(0.61-1.14)
Design:	Acutely ill	care	<b>G1b+G2b</b> : 140/676	<b>G2a</b> : 79/204 (30)	,
RCT	Undergoing	N at enrollment:	(20.7)	<b>G2b</b> : 126/338 (37)	Any test†‡, OR
	cancer	G1a+G2a: 498			(95%):
	treatment	<b>G1b +G2b</b> : 848	Any test†‡, n (%):	Any test, n (%):	G1a+G2a vs.
	Unresolved	N at followup:	<b>G1a+G2a</b> : 216/290	<b>G1a</b> : 117/146 (80)	<b>G2a+G2b:</b> 1.08
	abnormal	<b>G1a</b> : 239	(74.5)	<b>G1b</b> : 215/246 (87)	(0.76-1.54)
	screening	<b>G1b</b> : 431	<b>G1b+G2b</b> : 370/487	<b>G2a</b> : 111/144 (77)	
	Disparity	<b>G2a</b> : 259	(76.0)	<b>G2b</b> : 182/241 (76)	Most tests†‡, OR
	indicator(s):	<b>G2b</b> : 417	NA ( ( )   1   (0/)	<b>NA</b> (1) (0/)	(95%):
	Language	Length of	Most tests†‡, n (%):	Most tests, n (%):	G1a+G2a vs.
	barrier	followup:	<b>G1a+G2a</b> :133/290	<b>G1a</b> : 82/146 (56)	<b>G2a+G2b:</b> 1.37
	(Spanish-	18 months  Measure of	(45.9) <b>G1b+G2b</b> : 262/487	<b>G1b</b> : 181/246 (74) <b>G2a</b> : 66/144 (46)	(1.02-1.86), p<0.05
	speaking)			<b>G2b</b> : 129/241 (54)	All tootott OP
	Referent	fidelity: NR	(53.8)	GLD. 123/241 (04)	All tests†‡, OR (95%):
	group:	INIX	All tests†‡, n (%):	All tests, n (%):	G1a+G2a vs.
	Internal;		<b>G1a+G2a</b> :31/290	<b>G1a</b> : 42/146 (29)	G2a+G2b: 1.04
	English-		(10.7)	<b>G1b</b> : 98/246 (40)	(0.64-1.72)
	speaking		<b>G1b+G2b</b> : 54/487	<b>G2a</b> : 27/144 (19)	(0.0 / 1.1.2)
			(11.1)	<b>G2b</b> : 61/241 (25)	
			` /	` '	

Study	Baseline							
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity			
Beach et al., 2007 (continued)	Subgroup or secondary analysis description: Secondary analysis of data from an RCT of an intervention to improve screening among lowincome women. A subgroup analysis of relative efficacy of the intervention among Spanish versus English speaking participants				Disparity after intervention: Not explicitly reported. See outcome data.			

<sup>\*</sup> Up-to-date status: breast cancer (mammography within 18 months), cervical cancer (Pap test within 18 months), and colorectal cancer (home fecal occult blood test within 18 months).

<sup>†</sup>Excludes women with a hysterectomy (n=379)

<sup>‡</sup> Excludes women up-to-date at baseline on any screening test valid for ≥5 years (i.e. colonoscopy, barium enema, or sigmoidoscopy; n=276)

<sup>§</sup>Up-to-date status: breast cancer (mammography within 18 months); cervical cancer (Pap test within 18 months); and colorectal cancer (home fecal occult blood test within 18 months, a barium enema or sigmoidoscopy within 5 years, or a colonoscopy within 10 years). Patients with a hysterectomy or a baseline colorectal cancer screening test valid for 5 years or more were excluded from analyses (n = 569).

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Author: Coberley et al., 2007 Region/State: multistate, U.S. Setting: Members of 20 health plans and participants in diabetes disease management programs Enrollment period: NR Funding: NR Conflict of Interest: NR Design: Retrospective cohort	Condition: Diabetes mellitus Inclusion criteria:  • Members of Healthways health plans with diabetes identified by administrative claims  • At least 10 months of eligibility in Healthways diabetes disease management programs Exclusion criteria: NR Disparity: Race/ethnicity* Referent group: Internal; Non- health disparity zones, non- minority zip codes Subgroup or secondary analysis description: NA Measure of fidelity: NR	Quality improvement intervention(s): Diabetes disease management program (all patients), including telephone intervention to improve A1C testing rates in previously non-adherent members (lacked test in baseline period) Intervention target: Patients Groups: G1a: patients living in health disparity zone (HDZ), pre-intervention G1b: patients living in minority zip codes in HDZ, pre-intervention G1c: patients living in non-HDZ, pre-intervention G2a: patients living in HDZ, post-intervention G2b: patients living in non-minority zip codes in HDZ, post-intervention G2c: patients living in non-minority zip codes in HDZ, post-intervention G2C: patients living in non-HDZ, post-intervention M: 37,425 G1a: 3359 G1b: 2068 G1c: 34,066 G2a: 3359 G2b: 2068 G2c: 34,066 Length of followup: 12 months prior to start of disease management (baseline period) and 12 months following disease management intervention	NR Process: HbA1c testing rate, %: G1a: 51.8 G1c: 64.0	Clinical: NR Process: HbA1c testing rate, %: G2a: 59.4 G2c: 68.6 HbA1c testing rate relative increase†, % (p value): G1a: 7.6 (p<0.001) G2a: 4.6 (p<0.0001)	Disparity before intervention: Process: HbA1c testing rate, % G1a: 51.8 G2a: 64.0 Disparity after intervention: Process: HbA1c testing rate, %: G2a: 59.4 G2c: 68.6

<sup>\*</sup>Analyzed by whether members lived in minority zip codes (>50% of the population minorities) or health disparity zones (an area in which the diabetes disease prevalence was above the national average for minority zip codes).
†Difference from pre-intervention (i.e. baseline)

Additional baseline and outcome data reportedly only graphically

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study	addressing disp	parities in health o	Baseline	aucu)	
Description	Population	Intervention(s)	Characteristics	Outcomes	Dis parity
Author:	Condition:	Quality	Clinical:	Clinical:	Disparity before
Wells et al.,	Depression	improvement	PCS-12 score at	Mental Health	intervention:
2007; Wells et	Inclusion	intervention(s):	9 years, mean ±	Inventory, mean ±	NR
al., 2004;	criteria:	Partners in Care	SD:	SD:	Disparity after
Miranda et al.,	• Use the	(PIC) quality	Minority:	Displayed	intervention:
2004	practice for the		36.98 ± 10.95	graphically without	QI-ethnicity
Region/State:		programs included	White: 34.80 ±	numerical values	interaction,%
Across U.S.	Positive for	commitment of in-	10.24		depressed and not
Setting:	current	kind resources from	MCS-12 score at	Appropriate care	in appropriate care
Six	depressive	practices; trained	9 years, mean ±	at 6 month	at 57 months, t-test
nonacademic	symptoms and	clinician teams;	SD:	followup, %:	(p-value):
managed	probable	clinician toolkit and	Minority: 44.69 ±	QI-Meds, White:	QI-Meds: 0.67
Healthcare	depressive	patient education	10.81	35.2	(0.51)
Organizations;	disorder in the	materials; trained	White: 46.49 ±	QI-Meds, minority:	QI-Therapy: 1.83
46 clinics with	last year	nurse specialist;	11.55	36	(0.07)
181 primary	Exclusion	plus either	Process:		
care providers	criteria:	enhanced	NR	QI-Therapy, White:	Omnibus test of QI
Enrollment	<ul><li>&lt; 18 years old</li></ul>	medication		38.1	effects at 9 years,
period:	<ul> <li>Not fluent in</li> </ul>	management (QI-		QI-Therapy,	minority vs. White
June 1996 –	English or	Meds) or enhanced		minority: 45	(95% CI):
March 1997	Spanish	psychotherapy (QI-		110 14/1:	-11.40 (-27.97,
Funding:	<ul> <li>Lacked</li> </ul>	Therapy).		UC, White: 26.7	5.18)
National	insurance	Intervention		UC, minority: 19	p=0.176
Institute of Mental Health	coverage for	target: Patient		Depressed and not in appropriate	Interaction model† for disparity at 9
(NIMH)	the local	Groups:		care at 57 months,	
Conflict of	therapists	G1: QI-Meds		% (95% CI), p-	analyses
Interest:	participating in	G2: QI-Therapy		value for	estimate‡ (95%
NR	the	G3: Usual care		comparison vs.	CI), p value:
Design:	interventions Disparity:	(UC)		usual care:	QI-Meds, minority
RCT	Race/ethnicity	G1a: minorities, QI-		QI-Meds, White:	vs. White:
	(Minorities)	Meds		12.8 (5.3, 20.4)	-35.81 (-62.13,
	Referent group:	G1b: Whites, QI-		p=0.70	-9.50)
	Internal; White	Meds		QI-Meds, minority:	p=0.008
	Subgroup or	G2a: minorities, QI-		24.6 (16.5, 32.7)	
	secondary	Therapy		p=0.07	QI-Therapy,
	analysis	G2b: Whites, QI-			minority vs. White:
	description: NR	Therapy		QI-Therapy, White:	18.08 (-8.50, 44.66)
	•	G3a: minorities,		16.4 (10.6, 22.1)	p=0.179
		usual care		p=0.70	
		G3b: Whites, usual		QI-Therapy,	UC, minority vs.
		care		minority: 21.7 (13.1,	
				30.3)	(-47.36, 4.65)
				p=0.03	p=0.106
				LIC White: 44.7	
				UC, White: 14.7 (8.3, 21.1)	
				UC, minority: 34.3	
				(27.2, 41.5)	
				(21.2, 71.0)	

Study Baseline		
Description Population Intervention(s) Characteristics	Outcomes	Disparity
Wells et al., 2007; Wells et Minority mal., 2007; Wells et Minority minorit	nteraction nodel† for ntervention ffects at 9 years, djusted analyses stimate‡ (95%	Interaction model† for intervention group difference on disparity at 9 years, adjusted analyses estimate‡ (95% CI), p value: QI-Meds vs. UC: -14.46 (-49.47, 20.56) p=0.415 QI-Therapy vs. UC at 9 years: 39.44 (2.41, 76.47) p=0.037 QI-Therapy vs. Meds at 9 years: 53.90 (19.61, 88.18) p=0.002

<sup>†</sup>Interaction model (intervention status X ethnic minority)

<sup>‡</sup>Presented as area under curve over 9-year time period, derived from the 3-level mixed effects model.

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Dis parity
Author:	Condition: Diabetes mellitus Inclusion criteria: • African American and White New York City Medicare fee- for-service beneficiaries in senior centers, religious organizations, and senior housing complexes • Facilities and providers who serve this population were also targeted, including African- American serving physicians in New York City, identified participants of IPRO's physician office quality improvement project in the Bronx, Kings, New York and Queens counties, and African- American-	intervention(s): Five rounds of onsite visits with providers; initial visit consisted of introduction to IPRO Diabetes Disparities project, review of ADA guidelines for lipid disorder screening, and distribution of provider toolkits including decision support tools for diabetes prevention and management, provider reminder materials, and culturally and linguistically appropriate patient reminders and educational materials.  Additional visits provided technical assistance, promoted a CME	Clinical: NR Process: Receipt of biennial lipid profile, n (%): G1: 7,981 (63.8) G2: 12,313 (85.0) G3: 18,612 (82.8)	Clinical: NR Process: Receipt of biennial lipid profile, n (%): G1: 12,993 (80.5) G2: 13,942 (89.8) G3: 24,150 (89.7)	Disparity before intervention: Disparity in receipt of biennial lipid profile, %: G1 vs. G2: 21.2 G1 vs. G3: 19.0 Predictor of receipt of biennial lipid profile, OR (95% CI), p value: G1: 0.42 (0.37 to 0.47) G3: 1.00 p<0.0001 Disparity after intervention: Disparity in receipt of biennial lipid profile, %: G1 vs. G2: 9.3 G1 vs. G3: 9.2 Change in disparity in receipt of biennial lipid profile, %: G1 vs. G2: -11.9 G1 vs. G3: -9.8

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Mahotiere et al., 2006 (continued)	Exclusion criteria: Richmond County excluded due to small number of African American Medicare fee- for-service beneficiaries Disparity: Race/ethnicity (African American) Referent group: Internal; White Subgroup or secondary analysis description: NA Measure of fidelity: NR	interventions involved focus groups to understand characteristics, needs, and perspectives of AA patients. Culturally appropriate diabetes self-			

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Mahotiere et al., 2006 (continued)		Groups: G1: African- American Medicare fee-for-service (FFS) beneficiaries with diabetes residing in Bronx, Kings, New York and Queens counties. G2: White non-dual enrolled Medicare fee-for service beneficiaries in Bronx, Kings, New York and Queens counties G3: all White Medicare fee-for- service beneficiaries with diabetes in Bronx, Kings, New York and Queens counties N at enrollment: G1: 12,510 G2: 14,490 G3: 22,487 N at followup: G1: 16,140 G2: 15,526 G3: 26,923 Length of followup: Remeasurement was in 3/31/02- 3/31/04			

	addressing disp	arities in health o	Baseline	nucu)	
Study Description	Population	Intervention(s)	Characteristics	Outcomes	Dieparity
					Disparity
Author:	Condition:	Quality	Clinical:	Clinical:	Disparity before
Arean et al.,	Depression	improvement	HSCL-20	Response (>50%	intervention:
2005	Inclusion	intervention(s):		decrease in HSCL-	Any specialty
Region/State:	criteria:	Collaborative care	(range 0-4),	20 depression score	mental health visits
5 U.S. states	60 years or	model, including	mean ± SD:	from baseline),	or psychotherapy in
Setting:	older	stepped care	G1a+G2a:	adjusted %:	the past 3 months,
Primary care	Met criteria for	approach to	$1.7 \pm 0.6$	G1a: 42	n (%):
clinics	major	managing	G1b+G2b:	<b>G1b:</b> 54	<b>G1a+G2a</b> : 123 (9)
Enrollment	depression or	depression; primary		G1c: 42	<b>G1b+G2b</b> : 9 (4)
period:	dysthymia	care provider	G1c+G2c:	<b>G2a:</b> 19	<b>G1c+G2c</b> : 12 (9)
NR	according to	education;	$1.8 \pm 0.7$	<b>G2b:</b> 23	p=0.06
	SCID	depression clinical	Process:	<b>G2c:</b> 14	Disparity after
Funding:		specialist/	Any	G1a vs. G2a:	intervention:
John A. Hartford		depression care	antidepressant	p<0.001	Intervention effects
Foundation;	criteria:	manager works	use in the past 3	G1b vs. G2b:	on measures of
California	See inclusion	with patient and	months, n (%):	p<0.001	depression and
Healthcare	Disparity:	primary care	G1a+G2a:	G1c vs. G2c:	health-related
Foundation;	Race/ethnicity	provider,	594 (43)	p<0.001	functioning were of
Hogg	(Black and	medication	G1b+G2b:	D : : (U001 00	similar magnitude
Foundation;	Latino)	monitoring and brief		Remission (HSCL-20	•
Robert Wood		psychotherapy; use		depression score	among Whites.
Johnson	Referent group:	of clinical	62 (45)	<0.5), adjusted %:	
Foundation	Internal; White	information tracking		<b>G1a</b> : 24	No significant
	Subgroup or	system; ready	Any specialty	<b>G1b</b> : 33	interactions
Conflict of	secondary	access to a	mental health	G1c: 25	between
Interest:	analysis	psychiatrist	visits or	<b>G2a</b> : 8	intervention and
NR	description:	Intervention	psychotherapy in		ethnic group for
<b>.</b>	Randomization	target: Appropriate		<b>G2c:</b> 9	clinical outcomes,
Design:	not stratified by	management of	months, n (%):	G1a vs. G2a:	including measures
RCT (secondary		depression	G1a+G2a:	p<0.001	of depression and
analysis)	post hoc	Groups:	123 (9)	G1b vs. G2b:	health related
	subgroup	G1a: Collaborative	G1b+G2b:	p<0.001	functioning.
	analysis	care, White patients		G1c vs. G2c:	
	conducted to	G1b: Collaborative		p=0.014	
	compare	care, Black patients	12 (9)		
	outcomes in each	<b>G1c:</b> Collaborative	0 (	HSCL-20 depression	
	race/ethnic group	care, Latino	Satisfaction with	score (range 0-4),	
	between	patients	depression care	mean:	
	intervention and	<b>G2a:</b> Usual care,	(excellent/very	<b>G1a</b> : 1	
	control	White patients	good; only	<b>G1b</b> : 0.9	
		G2b: Usual care,	assessed in	G1c: 1	
		Black patients	those reporting	<b>G2a</b> : 1.39	
		<b>G2c:</b> Usual care,	depression care	<b>G2b</b> : 1.4	
		Latino patients	in past 3	<b>G2c:</b> 1.4	
			months), n (%):	G1a vs. G2a:	
		N at enrollment:	G1a+G2a:	p<0.001	
		<b>G1a+G2a</b> : 1388	226 (50)	G1b vs. G2b:	
		G1b+G2b: 222	G1b+G2b:	p<0.001	
		<b>G1c+G2c</b> : 138	52 (60)	G1c vs. G2c:	
		N at followup:			
		<b>G1a+G2a</b> : 1208			
		<b>G1b+G2b</b> : 186			
		<b>G1c+G2c:</b> 119			

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Arean et al.,		Length of	G1c+G2c:	p=0.002	
2005		followup: 12	21 (48)	Overall functional	
continued)		months		impairment (range 0-	
		Measure of		10), mean:	
		fidelity:		<b>G1a:</b> 3.6	
		NR		<b>G1b:</b> 3.7	
				<b>G1c:</b> 3.9	
				<b>G2a:</b> 4.5	
				<b>G2b:</b> 4.7	
				<b>G2c:</b> 4.7	
				G1a vs. G2a:	
				p<0.001	
				G1b vs. G2b:	
				p=0.005	
				G1c vs. G2c:	
				p=0.089	
				Process:	
				Any antidepressant	
				use, adjusted %:	
				<b>G1a</b> : 66	
				<b>G1b</b> : 62	
				G1c: 68	
				<b>G2a</b> : 50	
				<b>G2b</b> : 46	
				G2c: 44	
				G1a vs. G2a:	
				p<0.001	
				G1b vs. G2b:	
				p=0.036	
				G1c vs. G2c:	
				p=0.016	
				Any psychotherapy	
				or specialty mental	
				health visits,	
				adjusted %:	
				<b>G1a:</b> 55	
				<b>G1b</b> : 35	
				G1c: 42	
				<b>G2a:</b> 16	
				<b>G2b:</b> 14	
				<b>G2c:</b> 12	
				G1a vs. G2a:	
				p<0.001	
				G1b vs. G2b:	
				p=0.01	
				G1c vs. G2c:	
				p=0.005	

Study		·	Baseline	·	·
Description	<b>Population</b>	Intervention(s)	Characteristics	Outcomes	Disparity
Arean et al.,				Satisfaction	
2005				(excellent/very good)	
(continued)				with depression	
				care, adjusted %:	
				<b>G1a</b> : 76	
				<b>G1b:</b> 72	
				<b>G1c</b> : 71	
				<b>G2a:</b> 49	
				<b>G2b:</b> 42	
				<b>G2c:</b> 50	
				G1a vs. G2a:	
				p<0.001	
				G1b vs. G2b:	
				p=0.001	
				G1c vs. G2c:	
				p=0.035	

No disparity measured at outset of study; numbers for depression related baseline values look similar

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Dis parity
Author: Ferreira et al., 2005 Region/State: Illinois Setting: General medicine primary care outpatient firms at a Veterans Affairs Medical Center Enrollment period: May 2001- December 2002 Funding: Department of Veterans Affairs Health Services Research Division and National Institutes of Health grants Conflict of Interest: None Design: RCT	at the study medical center Exclusion criteria: • Personal or	intervention: Health care provider-directed intervention including education and feedback Intervention target: Appropriate screening for colorectal cancer Groups: G1: participants treated at firm randomized to intervention G1a: high literacy, intervention G1b: limited literacy, intervention G2a: high literacy, usual care G2b: limited literacy, usual care N at enrollment: G1a: 118 G1b: 79 G2a: 125 G2b: 60 N at followup*: G1a: 118 G1b: 79 G2a: 125 G2b: 60 Length of followup: 6 - 18 months after	Clinical: NA Process: Eligible participants did not have a FOBT in the previous year or FS/COL in the previous 5 years.	Clinical: NA Process: Colorectal cancer screening recommended: G1a: NR G1b: NR G2a: NR G2b: NR Colorectal cancer screen test completed: FOBT only (%): G1a: 21.2 G1b: 30.4 G2a: 15.2 G2b: 6.7 FS/COL only (%): G1a: 12.7 G1b: 17.7 G2a: 16.8 G2b: 21.7 Both FOBT and FS/COL (%): G1a: 5.1 G1b: 7.6 G2a: 4.0 G2b: 1.7 Any screening (%): G1a: 39.0 G1b: 55.7 G2a: 36.0 G2b: 30.0 G1a vs. G1b: p=0.35 G1b vs. G2b: p=0.002	Disparity before intervention: NA; all eligible participants did not have a FOBT in the previous year, or FS/COL in the previous 5 years. Disparity after intervention: No significant difference in overall CRC screening incidence between intervention and usual care among patients with high literacy. Individuals with limited literacy in the intervention group were more likely to complete any CRC screening tests than individuals with limited literacy in the intervention group.

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Ferreira et al., 2005 (continued)	Subgroup or secondary analysis description: Exploratory analysis of individuals with limited (<9 <sup>th</sup> grade level) versus high literacy (≥ 9 <sup>th</sup> grade level) as measured by the REALM instrument	Measure of fidelity:  • 15 of the 60 providers in the intervention firm did not participate in the initial workshop.  • 84% of the physicians and nurse practitioners attended at least one of the four feedback sessions.			

<sup>\*</sup>N at baseline is derived from the N at followup.

N at baseline indicates 185 in the control firm completed health literacy assessment, and 197 in the intervention firm completed health literacy assessment. Authors state that data was available for all patients included in the study.

Patients who died during followup are not included in the final analysis

Randomization was at the outpatient clinic level (one intervention clinic and one usual care clinic)

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study	addiessing disp	arities in health o	Baseline	na ca y	
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Author:	Condition:	Quality	Clinical:	Clinical:	Disparity before
Sherbourne et	Depression	improvement	MCS-12, mean:	Probable unmet	intervention:
al., 2004	Inclusion	intervention(s):	Women: 34.0	need for	NR Biographics of the
Region/State:	criteria:	QI- MEDS, QI-	Men: 36.39	appropriate care at 24 months, %*:	intervention:
Across U.S. Setting:		THERAPY; in-kind resources, local	PCS-12, mean: Women: 45.06	QI-Meds, women:	Significant main
Six		expert and staff	Men: 44.66	53	effect for gender:
nonacademic	year	nurses training	CESD, mean:	QI-Meds, men: 52	t=3.97, p=0.0001
managed	Older than 17	Intervention	Women: 45.59	QI-Therapy, women:	, , , , , , , , , , , , , , , , , , , ,
Healthcare	<ul> <li>No acute</li> </ul>	target:	Men: 43.09	54	Probable unmet
Organizations	medical	Increase		QI-Therapy, men: 57	
Enrollment	emergency	access/adherence		UC, women: 45	care, 3 way
period:	<ul> <li>Spoke English</li> </ul>	with any form of		UC, men: 57	interaction model
June 1996- March 1997	or Spanish	appropriate care for depression;		UC vs. QI-Meds, women: p=0.13	(sex X time X intervention):
Funding:	Insurance or	Improved health		UC vs. QI-Therapy,	F=2.16, p=0.028
AHRQ, National	public-pay arrangement to	and employment		women: p=0.10	1 –2.10, p–0.020
Institute of	cover	outcomes among		QI-Meds vs. QI-	Mental HRQOL, 3
Mental Health,	intervention	ethnic minorities.		Therapy, women:	way interaction
John D. and	care	Groups:		p=0.92	model (sex X time X
Catherine T.	Exclusion	G1: QI-Meds		UC vs. QI-Meds,	intervention):
MacArthur	criteria:	<b>G2:</b> QI-Therapy		men: p=0.53	F=2.12, p=0.031
Foundation Conflict of	See inclusion	<b>G3:</b> Usual care (UC)		UC vs. QI-Therapy, men: p=0.96	Employment status,
Interest:	Disparity:	N at enrollment:		QI-Meds vs. QI-	3 way interaction
NR	Sex Referent group:	1358		Therapy, men:	model (sex X time X
Design:	Internal	N at followup		p=0.59	intervention):
RCT	Subgroup or	(completed > 1			F=1.99, p=0.044
	secondary	followup		Mental HRQOL at	
	analysis	questionnaire):		24 months, mean:	
	description: NR	Women: 941		QI-Meds, women:	
	Measure of	Men: 358 N at 24 months:		40.18 QI-Meds, men: 40.25	
	fidelity:	NR		QI-Therapy, women:	
	NR	Length of		41.10	
		followup:		QI-Therapy, men:	
		24 months		41.41	
				UC, women: 39.22	
				UC, men: 38.47	
				UC vs. QI-Meds,	
				women: p=0.41 UC vs. QI-Therapy,	
				women: p=0.01	
				QI-Meds vs. QI-	
				Therapy, women:	
				p=0.56	

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Sherbourne et al., 2004 (continued)				UC vs. Ql-Meds, men: p=0.32 UC vs. Ql-Therapy, men: p=0.12 Ql-Meds vs. Ql- Therapy, men: p=0.55  Working at 24 months, %*: Ql-Meds, women: 65 Ql-Meds, women: 53 Ql-Therapy, women: 53 Ql-Therapy, women: 59 UC, women: 60 UC, men: 56 UC vs. Ql-Meds, women: p=0.23 UC vs. Ql-Therapy, women: p=0.10 Ql-Meds vs. Ql- Therapy, women: p=0.00 UC vs. Ql-Meds, men: p=0.00 UC vs. Ql-Meds, men: p=0.90 UC vs. Ql-Therapy, men: p=0.61 Ql-Meds vs. Ql- Therapy, men: p=0.61 Ql-Meds vs. Ql- Therapy, men: p=0.71	
				Process:	
				NR	

<sup>\*</sup>Percent derived from proportion

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study	addressing disp	parities in health o	Baseline	nucu)	
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Author:	Condition:	Quality	Clinical:	Clinical:	Disparity before
					intervention:
Miranda et al.,	Depression Inclusion	improvement	MCS-12, mean ±		
2003	criteria:	intervention(s): QI- MEDS, QI-	SD 610.630.360	Appropriate care %, (95% CI)	Disparity after intervention:
Region/State: Across U.S.			<b>G1a+G2a:</b> 36.9 11.1	` '	
Setting:		THERAPY; in-kind	<b>G1b+G2b:</b> 38.3	<b>G1a:</b> 30.0 (23.9-36.2) <b>G2a:</b> 21.8 (12.9-30.7)	No significant interaction
Six		resources, local	± 11.0	p=0.13	
nonacademic		expert and staff nurses training		<b>G1b:</b> 43.1 (33.0-53.2)	between intervention and
managed	year	Intervention	10.3	<b>G2b:</b> 28.8 (19.1-38.5)	
Healthcare	Older than 17	target:	PCS-12, mean ±		the rate of
Organizations	No acute	Increase	SD	<b>G1c:</b> 47.7 (41.6-53.8)	appropriate care or
Enrollment	medical	access/adherence		<b>G2c:</b> 38.9 (32.1-45.7)	work status
period:	emergency	with any form of	11.2	p=0.05	(Latinos and
June 1996-	Spoke English     Spoke English	appropriate care for		12 months,	African Americans
March 1997	or Spanish	depression;	± 9.9	Appropriate care %,	combined and
Funding:	Insurance or	Improved health	G1c+G2c: 45.4 ±		compared with
AHRQ, National	public-pay	and employment	12.0	<b>G1a:</b> 39.4 (31.7-47.1)	White patients).
Institute of	arrangement to	outcomes among	Process:	<b>G2a:</b> 26.4 (17.3-35.5)	Willia palionio).
Mental Health,	cover	ethnic minorities.	Appropriate	p=0.03	Significantly more
John D. and	intervention	Groups:	care, N* (%)	<b>G1b:</b> 55.7 (27.1-84.4)	improvement
Catherine T.	care Exclusion	G1a: Latino,	<b>G1a+G2a</b> : 51	<b>G2b:</b> 35.2 (7.8-62.6)	under
MacArthur	criteria:	intervention	(12.8)	p=0.33	interventions
Foundation	See inclusion	G1b: African	G1b+G2b: 28	<b>G1c:</b> 62.1 (57.2-67.1)	among the two
Conflict of	criteria	American,	(29.4)	<b>G2c:</b> 53.7 (45.7-61.7)	minority groups as
Interest:	Disparity:	intervention	G1c+G2c: 275	p=0.07	compared with
NR	Race/ethnicity	G1c: White,	(35.3)	Process:	White patients
Design:	(African	intervention	G1a+G2a vs.	6 months, Working,	(p=0.02) at 6
RCT	American, Latino)	G2a: Latino, usual	G1c+G2c,	% (95% CI)	months.
	Referent group:	care	p<0.001	<b>G1a:</b> 59.2 (52.4-66.1)	
	Internal; Non-	G2b: African	Employment	<b>G2a:</b> 60.0 (52.5-67.6)	
	minority	American, usual	status, N* (%)	p=0.80	
	Subgroup or	care	<b>G1a+G2a</b> : 243	<b>G1b:</b> 64.7 (52.3-77.0)	
	secondary	G2c: White, usual	(60.9)	<b>G2b:</b> 66.1 (54.7-77.5)	
	analysis	care	<b>G1b+G2b</b> : 69	p=0.84	
	description: NR	N at enrollment:	(42.3)	<b>G1c:</b> 68.0 (60.8-75.3)	
	Measure of	<b>G1a+G2a</b> : 398	<b>G1c+G2c</b> : 502	<b>G2c:</b> 61.5 (53.7-69.2)	
	fidelity:	<b>G1b+G2b</b> : 93	(64.5)	p=0.01	
	NR	<b>G1c+G2c</b> : 778			
		N at followup:			
		6 month:			
		All: 1150-1156			
		12 month:			
		All: 1075-1126			
		Length of			
		followup:			
		Self-administered			
		mail surveys:			
		baseline and every			
		six months for two			
		years			

Study			Baseline		
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Miranda et al.,				12 months, Working,	
2003				% (95% CI)	
(continued)				<b>G1a:</b> 60.1 (51.3-68.8)	
				<b>G2a:</b> 56.6 (47.1-65.9)	
				p=0.38	
				<b>G1b:</b> 70.3 (56.2-84.4)	
				<b>G2b:</b> 75.6 (67.1-84.0)	
				p=0.43	
				<b>G1c:</b> 66.0 (58.1-73.8)	
				<b>G2c:</b> 59.8 (51.6-68.0)	
				p=0.02	
				Counseling or	
				antidepressant use	
				at appropriate dose,	
				controls vs.	
				intervention, 6	
				months,%: G1a+G1b+G1c: 50.9	
				G2a+G2b+G2c: 39.7	
				p<0.001	
				Counseling or	
				antidepressant use	
				at appropriate dose,	
				controls vs.	
				intervention, 12	
				months,%:	
				G1a+G1b+G1c: NR	
				G2a+G2b+G2c:	
				NR	
				p=0.006	
				Ethnicity x	
				Intervention	
				interaction;	
				p=0.02	

<sup>\*</sup>N calculated

N varies for appropriate care and working data in outcome (1150-1156, 6 months) and (1075-1126, 12 months). The N for each ethnicity is not broken down further, hence percentages are used.

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Author: Barr et al., 2001 Region/State: Northeast U.S. Setting: Large group- model HMO Enrollment period: October 1995* Funding: CDC Conflict of Interest: NR Design: RCT	Condition: Breast cancer, screening Inclusion criteria:  • Women aged 50-75 years  • Bilateral mammography in 1 <sup>st</sup> quarter 1994 with no subsequent mammogram 18-21 months after initial mammogram  • Continuously enrolled in the health plan from April 1, 1994 through March 31, 1996 Exclusion criteria: See inclusion criteria Disparity: Insurance status (Medicaid or Medicare) Referent group: Internal/Commer cial insurance Subgroup or secondary analysis description: analysis of outcomes within each insurance status by intervention group; analysis of overall likelihood of followup mammography by insurance status	G2a: commercially insured, telephone reminder G2b: Medicare, telephone reminder G3: usual care G3a: commercially insured, usual care G3b: Medicare,	<b>G1b</b> : 0 <b>G2a</b> : 0 <b>G2b</b> : 0 <b>G3a</b> : 0	Clinical: NA Process: Subsequent bilateral mammogram, %: G1a: 44.4 G1b: 43.6 G2a: 55.6 G2b: 56.4 G3a: 39.4 G3b: 42.7 G1a vs. G2a vs. G3a: p=0.001 G1b vs. G2b vs. G3b: p=0.01	Disparity before the intervention: NR Disparity after the intervention: Estimated relative risk of subsequent mammography , RR (95%CI): G1b+G2b+G3b vs. G1a+G2a+G3a: 1.04 (0.93-1.14).

Study Description	Population	Intervention(s)	Baseline Characteristics	Outcomes	Disparity
Barr et al., 2001 (continued)	1 opulation	Length of followup: 5 months† N at followup: G1: 630 G1a: 448 G1b: 158 G2: 653 G2a: 437 G2b: 195 G3: 625 G3a: 421 G3b: 180 Measure of fidelity: 79% of women randomized to telephone reminders were	Citaracteristics	Outcomes	<b>Бізрапц</b>
		reached by phone			

Medicaid groups were all < 50 patients, and were excluded from the table.

<sup>\*</sup>The sample was defined and randomized initially in October 1995, though eligibility was assessed from the 1<sup>st</sup> quarter of 1994. † Followup began 10/24/1995 and ended 3/30/1996, two years after the last initial mammogram ( 3/30/1994). ‡ Per the sample definition, none had a subsequent mammogram at the start of the followup period

Evidence Table I-1. Characteristics and outcomes from studies of quality improvement interventions addressing disparities in health outcomes (continued)

Study	addressing disp	artics in nearth	Baseline	nucu)	
Description	Population	Intervention(s)	Characteristics	Outcomes	Disparity
Author:	Condition:	Quality	Clinical:	Clinical:	Change in disparity:
Connett et al.,	Coronary heart	improvement	Diastolic blood	Diastolic blood	Diastolic blood
1984	disease,	intervention(s):	pressure, mean	pressure, mean	pressure, % change
Region/State:	prevention	MRFIT	mm Hg ± SD:	mm Hg:	in mean mm Hg:
18 U.S. cities	Inclusion	multifactorial	<b>G1a+G2a:</b> 90.7 ±	<b>G1a:</b> 80.4	<b>G1a:</b> -11.4
Setting:	criteria:	intervention (Blood		<b>G1b</b> : 81.3	<b>G1b:</b> -13.6
Clinical centers	<ul> <li>Men, ages 35-</li> </ul>	pressure	<b>G1b+G2b</b> : 94.0 ±	<b>G2a:</b> 83.5	<b>G2a:</b> -7.9
Enrollment	57	management,	9.2	<b>G2b:</b> 85.6	<b>G2b:</b> -8.1
period:	<ul> <li>Elevated CHD</li> </ul>	dietary counseling,			
1973-1975	risk	smoking cessation	Systolic blood	Systolic blood	Systolic blood
Funding:	(Framingham	counseling)	pressure, mean	pressure, mean	pressure, % change
National Heart,	score in top 10-	Usual care	mm Hg ± SD:	mm Hg:	in mean mm Hg:
Lung and Blood	15%)	Intervention	<b>G1a+G2a:</b> 135.4	<b>G1a:</b> 121.4	<b>G1a:</b> -10.3
Institute	Exclusion	target:	[14.1]	<b>G1b</b> : 122.3	<b>G1b:</b> -11.7
Conflict of	criteria:	CHD risk factor	<b>G1b+G2b:</b> 138.5	<b>G2a:</b> 126.5	<b>G2a:</b> -6.4
Interest:	<ul> <li>Known CHD,</li> </ul>	control	± 14.9	<b>G2b</b> : 129.4	<b>G2b:</b> -5.8
NR .	diabetes,	Groups:			
Design:	diastolic blood	G1: MRFIT	Smokers, % ± SD:		Smokers, % change:
RCT	pressure ≥ 115,		<b>G1a+G2a:</b> 63.0 ±		<b>G1a:</b> -46.2
	total cholesterol		48.2	<b>G1b:</b> 38.0	<b>G1b:</b> -43.0
	≥ 350, use of	intervention	<b>G1b+G2b:</b> 68.7 ±		<b>G2a:</b> -29.0
	lipid-lowering	G1b: Black,	46.4	<b>G2b:</b> 42.9	<b>G2b:</b> -22.5
	drugs, body	intervention <b>G2:</b> usual care	Serum cholesterol	Serum cholesterol.	Corum abalactoral 9/
	weight ≥ 150%	<b>G2a:</b> White, usual		mean mg/dl:	Serum cholesterol, % change in mean
	desirable,	care	visit), mean mg/dl		mg/dl:
	dietary	<b>G2b:</b> Black, usual		<b>G1b</b> : 231.0	<b>G1a:</b> -7.3
	restrictions	care	<b>G1a+G2a:</b> 254.2 ±		<b>G1b:</b> -6.0
	incompatible with study	N at enrollment:	36.4	<b>G2b:</b> 237.3	<b>G2a:</b> -5.6
	underlying	<b>G1:</b> 6,428	<b>G1b+G2b:</b> 245.7		<b>G2b: -</b> 3.7
	condition likely	G1a: NR	± 37.8	Plasma	
	to interfere with	G1b: NR		cholesterol, mean	Plasma cholesterol,
	study	<b>G2:</b> 6,438	Plasma	mg/dl:	% change in mean
	participation	G2a: NR	cholesterol (2 <sup>nd</sup>	<b>G1a:</b> 228.5	mg/dl:
	Disparity	G2b: NR	screening visit),	<b>G1b</b> : 224.4	<b>G1a:</b> -6.4
	indicator(s):	<b>G1a+G2a:</b> 11,935	mean mg/dl ± SD:	<b>G2a:</b> 233.3	<b>G1b:</b> -6.5
	Race/Ethnicity	<b>G1b+G2b</b> : 931	<b>G1a+G2a:</b> 240.8 ±	<b>G2b</b> : 230.2	<b>G2a:</b> -4.5
	(Black)	N at 6 <sup>th</sup> annual	36.7		<b>G2b: -</b> 4.3
	Referent group:	followup:	<b>G1b+G2b</b> : 236.4	Plasma HDL	
	Internal; White	<b>G1</b> : 5754	± 38.4	cholesterol, mean	Plasma HDL
	•	<b>G1a</b> : 5338	DI 1:5:	mg/dl:	cholesterol, %
		<b>G1b</b> : 416	Plasma HDL	<b>G1a:</b> 41.3	change in mean
		<b>G2</b> : 5638	cholesterol,	<b>G1b</b> : 46.0	mg/dl
		<b>G2a</b> : 5227	mean mg/dl ± SD:		G1a: -2.6
		<b>G2b</b> : 411	<b>G1a+G2a:</b> 41.6 ±	<b>G2b:</b> 47.6	<b>G1b:</b> -6.5
		<b>G1a+G2a</b> : 10,565 <b>G1b+G2b</b> : 827	11.2 <b>G1b+G2b:</b> 48.8 ±		<b>G2a:</b> -2.6 <b>G2b:</b> -4.8
		G10+G20: 02/			<b>GZD4</b> .0
			15.8		

Description Population Intervention(s)	Characteristics	Outcomes	Disparity
Connett et al., 1984 (continued)  Subgroup or secondary analysis of Black and White* participants from the Multiple Risk Factor Intervention Trial (MRFIT) to identify intervention effects for reducing CHD risk factors  Connett et al., Subgroup or secondary followup: 6-8 years, average length of followup 7 years Measure of fidelity: NR	Plasma LDL cholesterol, mean mg/dl ± SD: G1a+G2a: 160.1 ± 35.8 G1b+G2b: 158.8 ± 39.1  Plasma triglycerides, mean mg/dl ± SD: G1a+G2a: 198.2 ± 147.2 G1b+G2b: 143.9 ± 97.0  Weight, mean lb ± SD: G1a+G2a: 189.1 ± 27.1	Plasma LDL cholesterol, mean mg/dl: G1a: 148.7 G1b: 148.3 G2a: 152.8 G2b: 153.8  Plasma triglycerides, mean mg/dl: G1a: 200.9 G1b: 153.9 G2a: 203.0 G2b: 147.1  Weight, mean lb: G1a: 187.7 G1b: 192.3 G2a: 190.1	Dis parity  Plasma LDL cholesterol, % change in mean mg/dl: G1a: -8.6 G1b: -8.5 G2a: -6.4 G2b: -5.4  Plasma triglycerides, % change in mean mg/dl: G1a: +1.7 G1b: +7.2 G2a: +3.3 G2b: +4.0  Weight, percent change in mean lb: G1a: -0.7 G1b: -0.2 G2a: +0.7 G2b: +0.7

<sup>\*</sup>White group includes 376 participants (2.9%) identified as an ethnic/racial group other than black or white

## **Appendix J. Strength of Evidence for Outcomes**

Table J1. Strength of evidence for outcomes in studies addressing disparities associated with race or ethnicity

race or ethnicity						
Outcome/ Intervention	Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
Colorectal cancer screening						
Multifaceted patient education/self management with limited language concordance and patient reminders	1 <sup>1</sup> (465) RCT	High	Unknown	Indirect	Imprecise	Insufficient
Depression Severity Score	e (HSCL-20)					
Collaborative care model plus clinical information tracking	1 <sup>2</sup> (1748) RCT	High	Inconsistent	Direct	Imprecise	Insufficient
Mental Health Inventory						
Collaborative care with medication adherance support.	1 <sup>3</sup> (474) RCT	High	Inconsistent	Direct	Imprecise	Insufficient
Collaborative care with cognitive behavioral therapy	1 <sup>3</sup> (474) RCT	High	Inconsistent	Direct	Imprecise	Insufficient
Depression symptoms						
Collaborative care model	1 <sup>4</sup> (396) RCT	Moderate	Inconsistent	Direct	Imprecise	Insufficient
Probable depressive disor	rder					
Collaborative care with medication adherance support.	2 <sup>5-7</sup> (1269) RCT	High	Inconsistent	Direct	Imprecise	Insufficient
Collaborative care with cognitive behavioral therapy	2 <sup>5-7</sup> (1269) RCT	High	Inconsistent	Direct	Imprecise	Insufficient
Use of antidepressant me	dications					
Collaborative care model plus clinical information tracking	1 <sup>2</sup> (1748) RCT	High	Inconsistent	Indirect	Imprecise	Insufficient
Collaborative care model	1 <sup>4</sup> (396) RCT	Moderate	Inconsistent	Indirect	Imprecise	Insufficient
HbA1c goal <7% achieved	d					
Patient education (cultural competency training)	1 <sup>8</sup> (3773) RCT	Low	Unknown	Indirect	Imprecise	Insufficient
BP goal achieved						
Patient education (cultural competency training)	1 <sup>8</sup> (3773) RCT	Low	Unknown	Indirect	Imprecise	Insufficient

Table J-1. Strength of evidence for outcomes in studies addressing disparities associated with race or ethnicity (continued)

race or eliminate (con	tillacaj					
Outcome/ Intervention	Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
LDL cholesterol level goal achieved						
Patient education (cultural competency training)	1 <sup>8</sup> (3773) RCT	Low	Unknown	Indirect	Imprecise	Insufficient
HbA1c test						
Patient reminder	1 <sup>9</sup> (37425) Cohort	High	Unknown	Indirect	Imprecise	Insufficient
Lipid test						
Multifaceted provider and community intervention	1 <sup>10</sup> (28650) Cohort	Moderate	Unknown	Indirect	Imprecise	Insufficient
Smoking cessation						
Patient education/self- management	1 <sup>11</sup> (11392) Cohort	Moderate	Unknown	Indirect	Imprecise	Insufficient
Frequency of PCI, CABG	, and cardiac cath	eterization			*	
Multifaceted patient and provider education intervention and AMI GAP impementation	1 <sup>12</sup> (2857) Cohort	High	Unknown	Indirect	Imprecise	Insufficient
Decreased systolic and d	iastolic blood pres	sure				
Patient education/self- management	1 <sup>13</sup> (636) RCT	Moderate	Unknown	Direct	Imprecise	Insufficient
Achieving diastolic blood	pressure goal					
Patient education/self- management	1 <sup>11</sup> (11392) Cohort	Moderate	Unknown	Direct	Imprecise	Insufficient

**Abbreviations:** AMI=acute myocardial infarction; BP=blood pressure; GAP=American College of Cardiology's AMI Guidelines Applied in Practice; HbA1c=glycosylated hemoglobin; LDL=low density lipoprotein; PCI=percutaneous coronary intervention; RCT=randomized controlled trial; SOE=strength of evidence

Table J2. Strength of evidence for outcomes in studies addressing disparities associated with insurance status

modiumoc otatas						
Outcome/ Intervention	Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
Colorectal cancer screening	ng					
Multifaceted patient education/self management with limited language concordance and patient reminders	1 <sup>1</sup> (465) RCT	High	Unknown	Indirect	Imprecise	Insufficient
Breast cancer screening						
Patient reminder system plus scheduling	1 <sup>14</sup> (3034) RCT	High	Unknown	Indirect	Precise	Insufficient

Table J3. Strength of evidence for outcomes in studies addressing disparities associated with

language

langaage						
Outcome/ Intervention	Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
Colorectal cancer screening	ng					
Language concordant patient education/self-management	1 <sup>15</sup> (1070) RCT	Moderate	Unknown	Indirect	Imprecise	Insufficient
Multifaceted patient education/self management with limited language concordance and patient reminders	1 <sup>1</sup> (465) RCT	High	Unknown	Indirect	Imprecise	Insufficient
Breast cancer screening						
Language concordant patient education/self-management	1 <sup>15</sup> (1346) RCT	Moderate	Unknown	Indirect	Imprecise	Insufficient

Table J4. Strength of evidence for outcomes in studies addressing disparities associated with health literacy

Outcome/ Intervention	Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
Colorectal cancer screening	•					
Multifaceted patient and provider education and reminder intervention with audit and feedback	1 <sup>16</sup> (1978) RCT	High	Unknown	Indirect	Imprecise	Insufficient

Table J5. Strength of evidence for outcomes in studies addressing disparities associated with socioeconomic status

~					
Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
edications					
1 <sup>17</sup> (1801) RCT	Moderate	Inconsistent	Indirect	Precise	Insufficient
1 <sup>4</sup> (396) RCT	Moderate	Inconsistent	Indirect	Imprecise	Insufficient
e (HSCL-20)				··	
1 <sup>17</sup> (1801) RCT	Moderate	Inconsistent	Direct	Precise	Insufficient
		•			
1 <sup>9</sup> (37425) Cohort	High	Unknown	Indirect	Imprecise	Insufficient
	Studies (N) Study Design  edications  1 <sup>17</sup> (1801) RCT  1 <sup>4</sup> (396) RCT  re (HSCL-20)  1 <sup>17</sup> (1801) RCT  1 <sup>9</sup> (37425)	Studies (N) Study Design  Pedications  1 <sup>17</sup> (1801) Moderate RCT  1 <sup>4</sup> (396) Moderate RCT  re (HSCL-20)  1 <sup>17</sup> (1801) Moderate RCT  1 <sup>9</sup> (37425) High	Studies (N) Study Design  Risk of Bias  Consistency  Edications  1 <sup>17</sup> (1801) Moderate Inconsistent  RCT  1 <sup>4</sup> (396) Moderate Inconsistent  RCT  Te (HSCL-20)  1 <sup>17</sup> (1801) Moderate Inconsistent  RCT  1 <sup>9</sup> (37425) High Unknown	Studies (N) Study Design  Risk of Bias  Consistency Directness  edications  1 <sup>17</sup> (1801) Moderate Inconsistent Indirect  1 <sup>4</sup> (396) Moderate Inconsistent Indirect  re (HSCL-20)  1 <sup>17</sup> (1801) Moderate Inconsistent Direct  RCT  1 <sup>9</sup> (37425) High Unknown Indirect	Studies (N) Study Design  Risk of Bias  Consistency Directness Precision  Precision  Precision  Risk of Bias  Consistency Directness Precision  Precision  Precise  Precise

**Abbreviations:** HbA1c=glycosylated hemoglobin; RCT=randomized controlled trial; SOE=strength of evidence

Table J6. Strength of evidence for outcomes in studies addressing disparities associated with sex

Outcome/ Intervention	Number of Studies (N) Study Design	Risk of Bias	Consistency	Directness	Precision	Overall SOE
Probable depressive disc	order					
Collaborative care with medication adherance support.	1 <sup>18</sup> (1299) RCT	Moderate	Inconsistent	Direct	Imprecise	Insufficient
Collaborative care with cognitive behavioral therapy	1 <sup>18</sup> (1299) RCT	Moderate	Inconsistent	Direct	Imprecise	Insufficient
Use of mental health serv	vices					
Collaborative care with medication adherance support.	1 <sup>18</sup> (1299) RCT	Moderate	Inconsistent	Indirect	Imprecise	Insufficient
Collaborative care with cognitive behavioral therapy	1 <sup>18</sup> (1299) RCT	Moderate	Inconsistent	Indirect	Imprecise	Insufficient

Table J7. Strength of evidence domains and domain definitions

Domain	Definition and elements	Score and application
Risk of bias	Risk of bias is the degree to which the included studies for a given outcome or comparison have a high likelihood of adequate protection against bias (i.e., good internal validity), assessed through two main elements:  • Study design (e.g., RCTs or observational studies)  • Aggregate quality of the studies under consideration. Information for this determination comes from the rating of quality (good/fair/poor) done for individual studies	Use one of three levels of aggregate risk of bias:  • Low risk of bias  • Medium risk of bias  • High risk of bias
Consistency	The principal definition of consistency is the degree to which reported effect sizes from included studies appear to have the same direction of effect. This can be assessed through two main elements:  • Effect sizes have the same sign (that is, are on the same side of "no effect")  • The range of effect sizes is narrow.	Use one of three levels of consistency:  Consistent (i.e., no inconsistency)  Inconsistent  Unknown or not applicable (e.g., single study)
Directness	The rating of directness relates to whether the evidence links the interventions directly to health outcomes. Evidence is indirect if:  • It uses intermediate or surrogate outcomes instead of health outcomes.  • It uses two or more bodies of evidence to compare interventions A and B.	Score dichotomously as one of two levels of directness:  • Direct • Indirect
Precision	Precision is the degree of certainty surrounding an effect estimate with respect to a given outcome (i.e., for each outcome separately).	Score dichotomously as one of two levels of precision:  • Precise • Imprecise

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## **Appendix K. Applicability Tables**

Table K-1. Applicability of evidence from cancer screening studies

Domain	Description of applicability of evidence
Population	Studies included patients cared for at community clinics in New York City, men treated at two Veterans Affairs clinics in Chicago, low-income primary care patients, men and women patients of a large urban primary care practice, and women enrolled in a large group model HMO in the northeastern US. As the populations were varied, study results may or may not be applicable to other areas.
Intervention	The studied interventions included provider education, audit and feedback, and patient education that was sensitive to low health literacy; mailed or telephone reminders to undergo screening; a targeted intervention by mail or mail intervention plus telephone reminders to encourage screening; patient navigator based education to undergo screening; and language-concordant patient education and care management. With the exception of mailed reminders (which were ineffective), these interventions required significant organizational resources to develop and implement and may not be feasible in other settings.
Comparators	The studies employed usual care as the comparator, which may differ in other settings.
Outcomes	Studies focused on process measures, namely, performance of breast, cervical, and colorectal cancer screening during follow-up. Included studies did not assess clinical outcomes, such as diagnosis of malignancies, leaving the long-term clinical impact unclear.
Setting	Studies were conducted in community clinics in New York City, VA clinics in Chicago, community health centers and public hospitals in Massachusetts, an academic primary care practice in Philadelphia, and a large group model HMO in the Northeast.

Table K-2. Applicability of evidence from cardiovascular disease studies

Domain	Description of applicability of evidence
Population	Studies of CAD risk factor control included men enrolled at clinical centers in 18 US cities and patients cared for at two university-affiliated primary care clinics in North Carolina. A study of AMI treatment included patients at academic and community hospitals in Michigan. The results of the latter two studies are applicable to ambulatory and hospitalized patients, respectively. However, the 18-city study, involving men only, has limited applicability to women.
Intervention	Interventions for CAD risk factor control included intensive patient education and self-management. The intervention for AMI treatment involved provider education, practice feedback, and a toolkit. The CAD risk factor interventions required significant institutional resources, and may or may not be feasible outside the research setting. The AMI intervention is part of a national guideline implementation initiative and has been applied in both research and clinical contexts.
Comparators	Usual care served as the comparator, which may differ in other settings.
Outcomes	Studies of CAD risk factor control reported intermediate clinical outcomes (blood pressure, cholesterol, smoking, weight). The study of AMI treatment assessed numerous process and utilization measures (prescription of evidence-based medications, smoking cessation counseling, use of the toolkit, echocardiography, cardiac catheterization, PCI, CABG), as well as proximal adverse clinical events (hypotension, shock, heart failure, pulmonary edema, stroke, renal insufficiency, bleeding). None of these studies reported subsequent health care utilization or longer-term clinical outcomes such as myocardial infarction (or reinfarction), or mortality, leaving the effect of the interventions on such issues unclear.
Setting	Among studies to improve CAD risk factor control, one was conducted in university-affiliated clinics; its results are applicable to academic primary care practices and may or may not be applicable to other outpatient primary care and subspecialty clinics. The other occurred in community clinics in 18 cities around the US, but enrolled patients between 1973 and 1975 and has limited applicability to present day practice. The AMI treatment intervention took place in academic and community hospitals, and is applicable to those settings.

Table K-3. Applicability of evidence from depression care studies

Domain	Description of applicability of evidence
Population	Two of the three studies focused on elderly patients in primary care. One included a range of ages in adulthood. All included both men and women and were racially diverse. Nonetheless, this represents a small proportion of the individuals who struggle with depression. It is unclear whether the observed results apply to apply to patient populations who receive their primary and mental health care outside of a managed care system, or to harder to reach individuals not receiving regular medical care. Given the settings in which the study took place, they also may not apply to vulnerable populations receiving care through public health systems.
Intervention	The intervention in all cases was some variation on a collaborative care model that included a care manager, usually a nurse. The interventions were all intensive in terms of demand on resources and required strong communication between care providers. One study had two interventions: one focused on providing training to the practice in medical management and one on therapy. Patients and their physicians, however, could also select any approach to managing their depression.
Comparators	All of the studies compared the intervention to usual care, although usual care was not ever completely described and therefore would be expected to vary.
Outcomes	Generally speaking, outcomes were appropriate and reflected those that would and could be used in practice. They included changes in depressive symptoms, incidence of probably depressive disorder, mental health related quality of life, functional impairment, and receipt of appropriate depression care.
Setting	All of the studies were conducted in primary care practices associated with larger healthcare organizations. It is unclear whether results would therefore apply to other settings, including individual practices without the resources of a larger organization, or assisted living facilities (pertinent because of the focus on the elderly population).

Table K-4. Applicability of evidence from diabetes care studies

Domain	Description of applicability of evidence
Population	Studies included people cared for by primary care clinicians in ambulatory health centers in eastern Massachusetts, diabetes disease management program members living in socioeconomic disparate areas throughout the U.S., and Medicare patients in New York State. Therefore, the results may or may not be applicable to other populations in other regions.
Intervention	Interventions evaluated included cultural competency training for clinicians and race-stratified performance reports with recommendations for Black diabetic patients, patient telephone reminders in Health Disparity Zones, defined as one with diabetes prevalence above the national average for minorities, and Medicare New York State Quality Improvement Organization (IPRO) multifaceted provider and community interventions. The interventions may not be available in other regions and settings, since they required significant programmatic and implementation resources.
Comparators	Comparators included usual care, process measures in Non-Health Disparity Zones; statewide interventions that IPRO implemented for the Physician Office Quality Improvement Project (POQIP). The usual care described in these studies may not be applicable to other settings and regions.
Outcomes	Studies reported surrogate clinical outcomes (HbA1c control), clinical risk factors for diabetes comorbidities (BP and lipid control), and process measures (HbA1c and LDL measurements). Duration of studies was generally 12 months. No studies reported any critically important clinical outcomes of diabetes such as death or micro-vascular and/or macro-vascular complications. Results from surrogate outcomes may not apply to important long-term clinical outcomes in people with diabetes.
Setting	Studies were conducted in ambulatory health centers in eastern Massachusetts, in diabetes disease management programs across the United States, and in New York state. As much of diabetes care is delivered in primary care ambulatory settings, the evidence would be applicable. However, specialty clinic settings were not reported and the evidence may not apply.

## **Appendix L. Excluded Papers**

Table L-1. Excluded papers: exclusion code, exclusion reason, and count

Exclusion code	Exclusion reason	Count
X-1	Not original research (e.g. review articles, systematic reviews, editorials, commentaries, letters to editor, etc.)	763
X-2	Does not include an intervention	2,880
X-3	Does not include individuals receiving health care within the U.S.	1, 123
X-4	The number of participants enrolled is less than 50 per group	2,758
X-5	Does not address a pre-specified condition*	669
X-6	Does not include individuals from a target population with selected characteristics known to be associated with health disparities†	1,937
X-7	Does not address a quality improvement strategy or intervention (i.e. a systematic process designed to improve the quality of care)	534
X-8	Intervention does not originate from or occur within a health care setting (i.e. hospital, clinic, or provider office)	59
X-9	Does not include a qualifying referent group‡	863
X-10	Does not addresses an outcome of interest (i.e. health outcome, process outcome, harm, or unanticipated adverse effect)	92
X-11	Does not include a control group for the intervention	19
X-12	Not a qualifying study design	8
X-13	Unable to obtain	7

## Notes:

\*Included: asthma; colorectal cancer (including screening); breast cancer (including screening); cardiovascular disease (including congestive heart failure, coronary artery disease, and hypertension); cystic fibrosis; depression (major depressive disorder only); diabetes mellitus; end stage renal disease; pregnancy; pneumonia (including pneumococcal vaccination); and pregnancy. †Included: race or ethnicity, socioeconomic status, insurance status, sex, sexual orientation, health literacy/numeracy, and language barrier.

‡A qualifying referent group was from either an internal (i.e., within study referent group) or external (i.e., data from a referent group not included in the study) source and had to provide adequate data (i.e. baseline characteristics and post-intervention outcome data) to facilitate an actual measure of disparity. Data from external referent groups needed to have been collected within four years of the enrollment of the target group and be from a source that was at the state or local level.

- 1. A study of women's awareness and use of mammograms. J Ky Med Assoc. 1987 Sep;85(9):553-5. PMID: 3668368. **X-2, X-4**
- 2. Hypoglycemia and employment/licensure. Diabetes Care. 1990 May;13(5):535. PMID: 2190777. **X-1, X-2, X-3, X-4, X-5, X-6**
- 3. Testimony of the American Dietetic Association: non-insulin-dependent diabetes mellitus--an unrelenting but undeserved threat to the health of minorities. J Am Diet Assoc. 1992 Jun;92(6):671-2. PMID: 1607556. **X-1**, **X-2**, **X-3**, **X-4**, **X-6**
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- 5. SSI (Supplemental Security Income) for premature infants. Continuum. 1995 Jul-Aug;15(4):17-8. PMID: 10145060. **X-1**, **X-2**, **X-3**, **X-4**, **X-5**, **X-6**
- 6. Setting the standards for asthma care. Bus Health. 1995 Jul;13(7 Suppl D):27-31. PMID: 10144994. **X-1**, **X-2**, **X-3**, **X-4**, **X-6**
- 7. A survey of treatment routines and educational level of health care providers in the initial phase of suspected acute myocardial infarction in Sweden in 1994. Swedish Working Group on Early Heart Attack Care. Eur J Emerg Med. 1996 Sep;3(3):149-56. PMID: 9023493. X-2, X-3, X-4, X-6
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- 9. Outcome data on Medicare patients with pneumonia. Connecticut Nursing News. 1997;70(9):10-1. **X-2**
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- 16. Quality compass '99: slow, but steady improvements. Med Manag Netw. 1999 Sep;7(9):5-7. PMID: 10558153. **X-1, X-2, X-3, X-4, X-5, X-6**
- 17. From the Centers for Disease Control and Prevention. Patients' reports of counseling on mammography screening by health-care providers--North Carolina, 1997. JAMA. 1999 Jul 14;282(2):124-5. PMID: 10411182. **X-6, X-7, X-9**
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- 19. Cutting RNs a false economy? Hosp Peer Rev. 1999 Feb;24(2):29-30. PMID: 10345889. **X-1, X-2, X-3, X-4, X-5, X-6**

- 20. Innovative breast cancer education programs for African-Americans. Oncology (Williston Park). 1999 Mar;13(3):298, 303. PMID: 10204152. **X-1**, **X-6**, **X-7**, **X-9**
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