Exercise Tolerance Testing to Screen for Coronary Heart Disease: A Systematic Review

By

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Abstract

Background: Coronary heart disease is the leading cause of morbidity and mortality in the United States. Exercise tolerance testing has been proposed as a means of better identifying asymptomatic patients at high risk for coronary heart disease events.

Purpose: To review the evidence on the use of exercise tolerance testing to screen adults with no history of cardiovascular disease for coronary heart disease. Data Sources: The MEDLINE database from 1966 through February 2003, handsearching of bibliographies, and expert input.

Study Selection: Eligible studies evaluated the benefits or harms of exercise tolerance testing when added to traditional risk assessment for adults with no known history of cardiovascular events.

Data Extraction: One reviewer extracted information from eligible articles into evidence tables, and another reviewer checked the tables. Disagreements were resolved by consensus.

Data Synthesis: No study has directly examined the effect of screening asymptomatic patients with exercise tolerance testing on coronary heart disease outcomes or risk-reducing behaviors or therapies. Multiple cohort studies demonstrate that screening exercise tolerance testing identifies a small proportion of asymptomatic persons (up to 2.7% of those screened) with severe coronary artery obstruction who may benefit from revascularization. Several large prospective cohort studies, conducted principally in middle-aged men, suggest that exercise tolerance testing can provide independent prognostic information

about the risk for future coronary heart disease events (relative risk with abnormal exercise tolerance testing, 2.0 to 5.0). However, when the risk for coronary heart disease events is low, most positive findings will be false and may result in unnecessary further testing or worry. The risk level at which the benefits of additional prognostic information outweigh the harms of false-positive results is unclear and requires further study.

Conclusions: Although screening exercise tolerance testing detects severe coronary artery obstruction in a small proportion of persons screened and can provide independent prognostic information about the risk for coronary heart disease events, the effect of this information on clinical management and disease outcomes in asymptomatic patients is unclear.

Introduction

Coronary heart disease is the leading cause of death in the United States. Each year, more than 1 million Americans experience nonfatal or fatal myocardial infarction or sudden death from coronary heart disease. Coronary heart disease can also present as angina, but only 20% of acute coronary events are preceded by long-standing angina (1). An estimated 1 to 2 million middle-aged men have asymptomatic but physiologically significant coronary artery obstruction, which puts them at increased risk for coronary heart disease events (2, 3). The economic burden of coronary heart disease is also substantial. The direct and indirect costs of coronary heart disease in the United States are projected to total \$129.9 billion for 2003 (1). The clinical and economic impact of coronary heart disease is the basis for considerable public health interest in the development of effective strategies to reduce the incidence of coronary heart disease events. In 1996, the U.S. Preventive Services Task Force considered the use of resting electrocardiography or exercise tolerance testing to detect asymptomatic coronary artery disease and prevent coronary heart disease events (4). The Task Force

found insufficient evidence to recommend for or against using these tests to screen middle-aged and older men and women. They recommended against screening children, adolescents, or young adults.

To update the evidence review and recommendations on screening for asymptomatic coronary artery disease, the Task Force and the Agency for Healthcare Research and Quality requested that the RTI International-University of North Carolina Evidence-based Practice Center perform an updated evidence

review beginning in 2001. The complete review considers resting electrocardiography, exercise tolerance testing, and electron-beam computed tomography for coronary calcium and is available at www.ahrq.gov. This article describes the findings on exercise tolerance testing only. The recommendations and rationale of the Task Force on screening for asymptomatic coronary artery disease are available at www.ahrq.gov (5, 6).

Clinicians can use 2 general approaches to prevention of morbidity and mortality from coronary heart disease. The first approach involves screening for and treating the traditional modifiable risk factors for coronary heart disease, such as hypertension, abnormal blood levels of lipids, diabetes, cigarette smoking, physical inactivity, and diet. Such an approach may incorporate explicit calculations of the patient's risk for coronary heart disease events by using risk prediction equations derived from the Framingham Heart Study or other cohort studies (7). The second strategy involves supplementation of screening based on traditional risk factors with additional tests to provide further information about future risk for coronary heart disease or to detect severe blockages of the coronary arteries that might warrant treatment.

Detection of increased risk for future coronary heart disease events may lead to intensified use of risk-reducing treatments. Some risk-reducing treatments are directed at traditional risk factors (for example, therapy with statins for hyperlipidemia), whereas others are not (for example, aspirin therapy). Revascularization by using coronary artery bypass graft surgery or percutaneous coronary intervention seeks to treat blockages of the coronary arteries. Whether

revascularization will reduce the risk for coronary heart disease events in persons identified by screening is unknown.

Exercise tolerance testing is widely used as a diagnostic test in the initial evaluation of patients with symptoms suggestive of myocardial ischemia and in persons with previously recognized coronary heart disease. Although exercise tolerance testing has been applied and studied as a screening or prognostic test in asymptomatic persons, its utility in this group is controversial. The best measure of the value of screening exercise tolerance testing would come from studies that examined whether patients randomly assigned to undergo such tests had fewer coronary heart disease events or received more appropriate risk-reducing therapies than did patients assigned to receive treatments after standard risk factor assessment.

Such direct evidence is not available. However, indirect evidence suggests that screening exercise tolerance testing may be helpful in guiding medical management (8). In the Multiple Risk Factor Intervention Trial Research study, high-risk male participants were randomly assigned to receive a multimodal intervention to reduce cardiovascular risk or usual care. Among participants with an abnormal baseline result on exercise tolerance testing, those who received the intervention had a significantly lower rate of mortality from coronary heart disease during follow-up than did the group that received usual care. No effect was seen among men with a normal baseline result on exercise tolerance testing. It is not clear from the report of this post-hoc analysis whether the cardiovascular

risk profiles of participants with an abnormal result on exercise tolerance testing at baseline differed significantly from those of participants with a normal result. Because direct evidence on possible benefits of screening exercise tolerance testing is lacking, we used data observational cohort studies to examine whether screening exercise tolerance testing could detect clinically significant asymptomatic obstructions of the coronary arteries or provide greater independent prognostic information about the risk for future coronary heart disease events than would be obtained solely by standard history, physical examination, and measurement of traditional risk factors. We also sought information about harms of screening, including the likelihood of false-positive results and the effect of labeling a person as being "at high risk."

Methods

Literature Review

To identify the relevant literature, we searched the MEDLINE database from 1966 through February 2003 by using the exploded Medical Subject Headings *coronary heart disease, exercise test,* and *mass screening* and the keywords *asymptomatic* and *screening*. We limited the search to English-language articles on human subjects. To supplement our literature searches, we hand-searched the bibliographies of key articles, used other recent systematic reviews when available, and included references provided by expert reviewers that had not been identified by other mechanisms.

Study Eligibility and Data Abstraction

Two reviewers examined the abstracts of the articles identified in the initial MEDLINE search and selected a subset for a full-text review. The same reviewers examined the full text of the selected articles to determine final eligibility. One reviewer extracted information from eligible articles into evidence tables, and another reviewer checked the tables. They resolved disagreements by consensus. To be eligible, studies had to have been performed in participants with no history of cardiovascular disease or to provide subset analysis for this group. Included studies on the detection of severe coronary artery obstruction reported the total number of persons screened to obtain the sample of persons with an abnormal result on exercise tolerance testing and the proportion of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number of participants found to have abnormal results on angiography by the total number screened.

For the prognostic benefit of exercise tolerance testing, included studies reported the independent value of the test for predicting coronary heart disease events. We included studies that examined the prognostic benefit of exercise testing by using several different variables, including ST-segment depression, functional capacity, chronotropic incompetence, heart rate recovery, and development of exerciseinduced premature ventricular contractions. We also included studies that used nuclear medicine imaging to detect ischemia. We excluded studies that did not use statistical methods to control for the effect of other risk factors (such as age or systolic blood pressure) on the estimate of the prognostic strength of a positive

result on exercise tolerance testing. Table 1 shows information on excluded studies.

The studies used different means of characterizing the prognostic benefit of screening with exercise tolerance testing. Many studies reported outcomes in terms of independent relative risk associated with a positive (versus a negative) screening test. Others used diagnostic test terminology, such as "sensitivity and specificity" or "positive predictive value." In such cases, the terms are used to indicate test accuracy over the entire follow-up period rather than at 1 point in time.

To assess whether a relationship exists between sensitivity of exercise tolerance testing for future coronary heart disease and duration of follow-up, we examined the correlation between reported sensitivity and mean duration of follow-up by using STATA statistical software, version 7.0 (Stata Corp., Chicago, Illinois).

Data Summary and Quality Assessment

We rated the quality of the included articles according to criteria developed by the U.S. Preventive Services Task Force Methods Work Group (9). Tables 3 and 4 show information only from studies judged "good." For the studies shown in Table 2, we considered several factors that affect quality, chiefly the percentage of patients with a positive exercise tolerance testing who underwent catheterization and how completely outcomes were assessed. We used the final set of eligible articles to create evidence tables and produce the larger evidence report, which also included evaluation of resting electrocardiography and electron-beam computed tomography to detect coronary calcium. The full

evidence report was subjected to external peer review and revised on the basis of the comments received; we used the revised report as the basis for this article. Role of the Funding Agency

This evidence report was funded through a contract to the RTI-University of North Carolina Evidence-based Practice Center from the Agency for Healthcare Research and Quality. Staff of the funding agency contributed to the study design, reviewed draft and final manuscripts, and made editing suggestions.

Results

We identified 713 articles for review. We reviewed the abstracts and retained 55 articles that examined the diagnostic or prognostic significance of screening with exercise tolerance testing. After full article review, we kept 31 articles representing 29 studies that met the inclusion criteria (10--40). We identified another 11 articles for inclusion through review of reference lists and input of expert reviewers (8, 41--50). Table 1 lists articles that were excluded during review of the full articles and the reason for exclusion (51--74).

We found no studies that directly tested whether screening asymptomatic persons with exercise tolerance testing improves coronary heart disease and mortality. Similarly, we found no studies that examined the effect of screening with exercise tolerance testing on the subsequent use of risk-reducing interventions and behaviors. However, we identified fair- or good-quality observational cohort studies of asymptomatic adults that prospectively evaluated the value of exercise tolerance testing in detecting asymptomatic coronary artery obstruction (14--18, 22, 23, 25, 27, 28, 30, 31, 38, 75) and predicting future coronary heart disease

events, such as angina, myocardial infarction, and sudden death (8, 10--13, 19--21, 26, 29, 32--36, 38--50). We also identified 3 good-quality studies that estimated the cost effectiveness of exercise tolerance testing to identify asymptomatic, severe, prevalent coronary heart disease (24, 28, 37). Exercise Tolerance Testing To Detect Asymptomatic Prevalent Disease We identified 13 studies in 14 articles that examined the utility of exercise tolerance testing to detect asymptomatic coronary artery obstruction (Table 2) (14, 15, 18, 22, 23, 25, 27, 28, 30, 31, 38, 75). In these studies, the prevalence of abnormal exercise tolerance testing, usually defined as exercise-induced STsegment depression of 1 mm or more, ranged from about 3% among aviators who were presumed healthy (16) to 29% in a sample of diabetic persons in Finland (15, 75). A portion of the participants with a positive exercise tolerance testing in each study (1% to 60%) proceeded to evaluation with cardiac catheterization. Screening with exercise tolerance testing yielded angiographically demonstrable coronary heart disease, usually defined as greater than 50% stenosis of a major coronary artery, in a minority of the screened patients.

The yield of screening exercise tolerance testing was greater in higher-risk groups. Five studies in 6 articles evaluated diabetic persons (15, 75), those with multiple risk factors (18, 31), those with siblings with coronary heart disease (17) and those who were prescreened by using a chest pain questionnaire (25). In these studies, the yield of screening for angiographically demonstrable coronary heart disease ranged from 1.2% (31) to 9% (15, 18). Most cases of coronary artery obstruction identified by screening were single-vessel disease, but up to 2.7% of

screened participants had significant left main or three-vessel disease (18) and as many as 1.7% proceeded to revascularization after screening (25). Eight studies screened unselected, low-risk patients (14, 16, 22, 23, 27, 28, 30, 38). These studies demonstrated a yield of 0.06% to 1.6% for asymptomatic coronary heart disease on angiography.

Cost Effectiveness

Three studies attempted to estimate the cost-effectiveness of screening to identify prevalent coronary artery obstruction. Sox and colleagues (24) used a decisionanalysis model to estimate the clinical effectiveness and cost-effectiveness of exercise testing in asymptomatic adults. Their model was structured so that the benefit of screening was achieved through detection of patients with severe disease who would benefit from revascularization. Only direct costs were considered. Levels were based on reimbursement rates at the time of the study (late 1980s): \$165 for exercise testing, \$3595 for angiography, and \$31 178 for coronary artery bypass surgery. No discounting rate was given. Screening 60-year-old men had a cost per life-year saved of \$24 600; for 60-year-old women, the cost was \$47 606. For persons 40 years of age, the cost-effectiveness ratios were much higher: \$80 349 per life-year saved for men and \$216 496 per life-year saved for women.

The presence or absence of risk factors for coronary heart disease affected the cost-effectiveness ratios. The cost per life-year saved was \$44 332 for 60-year-old men with no risk factors and \$20 504 for those with 1 or more risk factors. The investigators concluded that routine screening was not warranted in general but

that it may be beneficial for persons at increased risk for coronary heart disease (for example, older men with 1 or more risk factors). An earlier cost-effectiveness analysis of screening exercise tolerance testing had similar findings (37). Pilote and colleagues (28) performed a cost analysis of data from their study of the clinical yield of screening exercise tolerance testing to detect unsuspected severe coronary artery obstruction. They sampled more than 4000 persons referred to the Cleveland Clinic for screening exercise tolerance testing. Data on cost were obtained from 1994 Medicare reimbursement rates: \$110 for exercise testing, \$1780 for angiography, and \$27 270 for coronary artery bypass surgery. Screening identified 19 patients with severe coronary artery obstruction (0.44% of the cohort); of these, 14 had subsequent coronary artery bypass graft surgery. The investigators estimated a cost of \$39 623 to identify 1 case of severe coronary artery disease by screening exercise tolerance testing. The estimated cost per year of life saved was \$55 274.

On the basis of these studies, it appears that screening with exercise treadmill testing and performing bypass surgery on persons with severe obstructions is relatively cost effective compared with other, better-accepted types of preventive care, such as mammography in women 50 to 69 years of age (76).

Exercise Tolerance Testing as a Prediction Tool for Risk for Coronary Heart Disease Events

Exercise tolerance testing can be used to provide information about a person's risk for a future coronary heart disease event that may augment the predictive ability of traditional risk assessment. Better risk assessment may help clinicians

and patients make better decisions about interventions for intermediate- and longterm risk reduction.

ST-Segment Response

Traditionally, studies of the predictive value of exercise tolerance testing on future coronary heart disease have examined ST-segment response to exercise as the risk predictor. Most of these studies reported the total number of coronary heart disease events (fatal and nonfatal myocardial infarction, new-onset stable or unstable angina, and coronary death) as their main outcome. Others reported death from coronary heart disease or from all causes as the main outcome or as secondary outcomes. The mortality rate from coronary heart disease, and particularly the total mortality rate, may be less subject to ascertainment bias than is the total number of coronary heart disease events and hence may be more valid measures. However, whether from coronary heart disease or other causes, death is uncommon in the generally healthy, asymptomatic patients enrolled in these studies, making it difficult to estimate the ability of exercise tolerance testing to predict such events.

We identified 15 studies in 18 articles that examined the relationship between ST segment response to exercise and risk for future coronary heart disease events (Table 2) (8, 11--13, 19--21, 26, 29, 32, 33, 36, 39--42, 45, 50). Thirteen of these studies (in 16 articles) found that ST-segment response during exercise predicted future coronary heart disease events (8, 11--13, 19--21, 26, 29, 33, 36, 39--41, 45, 50). In 1 of these studies, only coronary heart disease events occurring during exercise was considered as the outcome (12); we therefore excluded it from

analysis of the predictive utility for coronary heart disease events . Two studies found that ST-segment response to exercise alone did not predict future coronary heart disease events (32, 42).

Of the studies that found ST-segment response to be predictive of future coronary heart disease events, 6 (published in 8 articles) selected persons for participation on the basis of the presence of 1 or more risk factors: diabetes (13), multiple risk factors (8, 33, 39, 50), hyperlipidemia (26, 41), and sedentary lifestyle and obesity (29). The prevalence of an abnormal exercise tolerance testing, usually defined as ST-segment depression of 1 mm or more, ranged from 12% to 52%. After adjustment for other risk factors, the independent relative risk for coronary heart disease events associated with an abnormal ST-segment response to exercise in these higher-risk groups ranged from 3.5 (8, 50) to 21.0 (13). Sensitivity for occurrence of coronary heart disease events over the duration of the studies (3 to 8 years) ranged from 30% to 100%. The positive predictive value of an abnormal exercise tolerance testing ranged from 7.1% (26, 41) to 46% (29).

Seven studies (published in 8 articles) found ST-segment response to exercise to be predictive of future coronary heart disease events in an unselected, low-risk sample (11, 19--21, 33, 36, 40, 45). The prevalence of an abnormal test tended to be lower than that in the higher-risk sample, ranging from 3% (33) to 20% (11, 21). The independent relative risk for coronary heart disease events associated with an abnormal exercise tolerance testing ranged from 1.6 (40) to 21 (33), with the majority of the values between 2.0 and 5.0. Gibbons and colleagues (33) reported a higher relative risk in low-risk persons (21.0) than did the other

investigators; however, the absolute event rate was low (0.08 to 2.8 events/1000 person-years) and the confidence interval was wide (6.9 to 63.3). The sensitivity of exercise tolerance testing for coronary heart disease events was 10% (45) to 70% (11, 21). The positive predictive values ranged from 2.2% (33) to 24% (19). Two of the studies added nuclear perfusion imaging to exercise electrocardiography (19, 32). These studies reported positive predictive values of about 50%. However, imaging is likely to increase screening program costs (19, 32).

As might be expected, the sensitivity of an abnormal exercise tolerance testing decreased as the duration of follow-up increased (r = -0.56). Data from these cohort studies suggest that the majority of asymptomatic persons with an abnormal exercise tolerance testing do not go on to have coronary heart disease events, at least within the time frame of follow-up. Persons who do have events often develop angina rather than experience myocardial infarction or sudden death. The prevalence of an abnormal result on exercise tolerance testing and its predictive value among asymptomatic persons is greater in those at higher risk. These data are consistent with those of other investigators and policymakers who have suggested that the value of exercise tolerance testing is greater when it is applied to patients with 1 or more risk factors for coronary heart disease because selection of a higher-risk cohort for screening increases the prevalence of disease and positive predictive value (10). Bruce and associates (10) reported that in the Seattle Heart Watch Study of 4158 asymptomatic men and women, a positive result on exercise tolerance testing in the absence of risk factors provided little

predictive value. However, among patients with 1 or more other risk factors for coronary heart disease, the occurrence of 2 different types of abnormal response to exercise tolerance testing (exercise risk predictors) was associated with a 15fold increase in risk compared with patients who had a normal result.

Other Exercise Predictors

More recent studies of the value of exercise testing in asymptomatic persons have examined the utility of other exercise-associated risk markers, including functional capacity, chronotropic incompetence, heart rate recovery, and development of exercise-induced premature ventricular contractions, for predicting patients' risk for coronary heart disease events or death (Table 3) (21, 34, 35, 42--49). In contrast to ST-segment response, these exercise indicators may not directly detect ischemic myocardium, but they probably indicate other cardiovascular derangements, such as abnormal autonomic regulation, that predict coronary heart disease events. In general, these findings are associated with moderate increases in risk for coronary heart disease after adjustment for other risk factors for coronary heart disease (relative risk, 1.7 to 3.5). Some factors are common: For example, failure to achieve target heart rate was noted in 21% of patients in the Framingham Offspring Study (44).

Exercise Tolerance Testing in Women

Two recent studies contribute important information on the predictive value of exercise tolerance testing in asymptomatic women (42, 43). The majority of other studies that we identified did not include women or did not provide subgroup analysis of the predictive value of screening exercise tolerance testing for women.

Mora and colleagues (42) analyzed data from the female participants in the Lipid Research Clinics Prevalence Study, many of whom had hyperlipidemia. They found that unlike in studies whose samples comprised predominantly men, STsegment response did not predict future risk for coronary heart disease events (relative risk, 0.88 [95% CI, 0.48 to 1.61]) in women (42). Low exercise capacity, along with low heart rate recovery after exercise, was an independent predictor of death from coronary heart disease (relative risk, 3.52 [95% CI, 1.57 to 7.86) and of all-cause death (relative risk, 2.11 [95% CI, 1.47 to 3.04]) in women. Gulati and coworkers (43) sampled asymptomatic female volunteers living in the Chicago area. They found that exercise capacity predicts risk for all-cause death in women. For every increase in exercise capacity of 1 metabolic equivalent, the relative risk for death was 0.83 (95% CI, 0.78 to 0.89). The predictive utility of exercise markers other than ST-segment response in these 2 studies of women is consistent with the results of similar studies in which most participants were men. Exercise Tolerance Testing before Beginning an Exercise Program Exercise tolerance testing is frequently used as part of an evaluation of middleaged persons before they begin an exercise program. Few data are available to determine the effectiveness of this approach in reducing the risk for activityrelated coronary heart disease events. Siscovick and colleagues (12) analyzed the effectiveness of exercise tolerance testing to predict activity-related coronary heart disease events in the Lipid Research Clinics cohort of asymptomatic hypercholesterolemic men. After an initial exercise tolerance test, the cohort was followed for an average of 7.4 years; during that time, the investigators used

retrospective record review to identify coronary heart disease events that were associated with moderate or intense activity. The cumulative incidence of activity-related coronary heart disease events during follow-up was 2%. An abnormal ST-segment response to exercise at the time of entry into the study was associated with a relative risk of 2.6 (95% CI, 1.3 to 5.2) for activity-related coronary heart disease events. The sensitivity of exercise testing for predicting the events was 18%, and the predictive value of a positive test for coronary heart disease events during exercise was 4%. Of the persons who had an activityassociated coronary heart disease event, 80% had an initially normal ST-segment response to exercise; 94% of persons with abnormal ST-segment response to exercise did not have an activity-associated event during follow-up. Thus, exercise testing appears to have limited ability to detect persons who will have exercise-related coronary heart disease events.

Adverse Effects of Screening Exercise Tolerance Testing Other than information on the frequency of false-positive results, we found no studies that examined the potential harms of screening. No study reported rates of complications from angiography of asymptomatic persons, measures of anxiety from knowledge of an abnormal test result, or adverse events from medical therapy initiated because of an abnormal test result.

Discussion

We identified no randomized trials that examined the effect of screening exercise tolerance testing to guide management and improve health outcomes of coronary heart disease or affect the use of risk-reducing treatments in asymptomatic adults.

Exercise tolerance testing of asymptomatic persons rarely detects previously unrecognized, clinically important coronary artery obstruction (up to 2.7% of screened persons). It does provide some independent prognostic information in at least some persons (relative risk of about 2.0 to 5.0 for coronary heart disease events associated with an abnormal result) above and beyond the prognostic information that can be gained from traditional assessment of risk factors. The effect of this additional information on clinical decision making, however, has not been studied. The potential benefits of screening exercise tolerance testing are likely to be small for groups in which the prevalence of the disease is low, such as young adults; such screening would also produce many cases of false-positive results. In such cases, the costs and harms associated with additional testing may exceed any benefits from screening.

The value of screening exercise tolerance testing rests in large part on the underlying incidence of coronary heart disease events and the prevalence of serious artery obstructions in the screened sample. Exercise tolerance testing will probably perform better when applied to higher-risk groups, such as persons with 1 or more risk factors coronary heart disease. Selection of a higher-risk group for screening increases the prevalence of disease in those screened and, thus, the predictive value of a positive test result. Whether the benefits of such tests exceed the disadvantages, including costs, in higher-risk groups is still unclear at present and requires investigation.

For persons at low risk for coronary heart disease events, a positive result on exercise tolerance testing is much more likely to be false positive than true

positive. False-positive results in this context are concerning because they can lead to unnecessary, and possibly injurious, additional procedures. Screening has been advocated for people with high-risk occupations, but we did not identify new studies on the effect of screening such patients. Data from studies of patients with known coronary heart disease but no ischemic symptoms suggest that treatment with medications, such as β -blockers, or revascularization can improve outcomes over no treatment, but whether patients with no history of coronary heart disease would have the same results is unclear (77). Exercise tolerance testing can be normal or nondiagnostic in an important proportion of patients who will experience a coronary heart disease event, as evidenced by the sensitivity values of 10% to 74% in the studies that evaluated ST-segment depression as a risk marker (Table 3). In a defined cohort of low-risk patients, a larger absolute number of coronary heart disease events occurs among those with an initially normal result on exercise tolerance testing than among those with an initially abnormal result. The suboptimal sensitivity of ST-segment response for predicting coronary heart disease events may be explained in part by the fact that ST-segment depression on exercise tolerance testing detects ischemia from obstructed coronary arteries, but many acute coronary heart disease events result from sudden occlusion of a previously nonobstructed segment of artery (78). Use of other measures from the exercise test that are not as dependent on identification of atherosclerotic obstructions may mitigate this dilemma (79). The primary tangible harm of screening exercise tolerance testing is the potential for medical complications related to cardiac catheterization done to further

evaluate a positive result. Coronary angiography is generally considered a safe procedure. Of all persons undergoing outpatient coronary angiography, however, an estimated 0.08% will die as a result of the procedure and 1.8% will experience a complication (80). Complications of coronary angiography include myocardial infarction, stroke, arrhythmia, dissection of the aorta and coronary artery, retroperitoneal bleeding, femoral artery aneurysm, renal dysfunction, and systemic infection. Rates of complications are likely to be somewhat lower in asymptomatic persons, but no good data are available. A positive result on exercise tolerance testing may also be an impetus to initiate risk-reducing therapy; hence, another potential harm of screening is use of with such therapies as aspirin or statins to overtreat persons who would not otherwise require treatment (that is, would be considered low risk) if they did not have an abnormal result on exercise tolerance testing. Other potential harms, including the psychological consequences of a false-positive test result, also have not been well studied. Our findings are consistent with those of the American Heart Association/American College of Cardiology expert panel, which also examined the effectiveness of screening exercise tolerance testing (33). They recommended against routine exercise tolerance testing in asymptomatic adults because of concerns about the positive and negative predictive value of screening exercise tolerance testing and the potential harms of false-positive results. The American Heart Association/American College of Cardiology found that screening exercise tolerance testing for persons with multiple risk factors to guide to risk-reduction

therapy or for sedentary middle-aged adults who wish to start a vigorous exercise program is controversial but potentially beneficial.

Further studies are required to determine the balance of benefits and harms of screening exercise tolerance testing for patients with different degrees of risk for coronary heart disease. An adequately powered randomized trial of screening exercise tolerance testing compared with management based on traditional risk factors would greatly inform clinical decision making. Such a study should compare a traditional global coronary heart disease risk assessment tool to a screening strategy that also incorporates exercise tolerance testing. A broad spectrum of patients should be enrolled, including a sufficient number of women. Studies examining how providers and patients actually apply the additional information from exercise tolerance testing will also be helpful. Finally, better information about the adverse effects of screening is required if researchers are to perform well-informed cost-effectiveness analyses of exercise tolerance testing screening plus risk factor-based decision making compared with risk-factor-based decision making alone.

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Table 1. Ex	cluded Studies	and Reasons	for Exclusions
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Author and Year	Reason for Exclusion
Allen et al., 1980 (51)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Aronow et al., 1975 (52;53)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Cumming et al., 1975 (54)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Elamin et al., 1982 (55)	Diagnostic usage – symptomatic patients
Fadayomi et al., 1987 (56)	Unclear ascertainment of endpoints
Froelicher et al., 1974 (57)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Froelicher et al., 1977 (58)	Did not report the total number of persons screened
Gerson et al., 1988 (59)	Did not report the independent risk of a positive exercise
	tolerance testing
<u>Gianrossi et al., 1989 (60)</u>	Diagnostic usage – symptomatic patients
<u>Goodman et al., 1989 (61)</u>	Subjects had previous history of cardiovascular disease
Gupta et al., 1983 (62)	Did not report the independent risk of a positive exercise
	tolerance testing
Hopkirk et al., 1984 (63)	Did not report the total number of persons screened
MacIntyre et al., 1981 (64)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Manca et al., 1982 (65)	Did not report the independent risk of a positive exercise
	tolerance testing
Mark et al., 1989 (66)	Subjects had previous history of cardiovascular disease
McHenry et al., 1984 (67)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Melin et al., 1981 (68)	Diagnostic usage – symptomatic patients
Pedersen et al., 1991 (69)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Roger et al., 1998 (70)	Included symptomatic patients without sub-analysis
Rubler et al., 1987 (71)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Selvester et al., 1996 (72)	Used a screening protocol that employed multiple
	technologies
l ubau et al., 1989 (73)	No adjustment of the relative risk of abnormal exercise
	tolerance testing for the effect of other risk factors
Uhl et al., 1981 (74)	Did not report the total number of persons screened

Author and Year	Study Population	Exclusion Criterion	Test Technique	Definition of Abnormal Exercise ECG
Caralis et al., 1979 (27)	3,496 men and women	NR	Maximal and thallium scintigram	≥2 mm of horizontal ST depression
	Mean age: NR % male NR		-	
Piepgrass et al., 1982 (16)	771 men in United States Air Force flight crew Mean age: (SD): 42 (5.2) 100% male	Rest ECG abnormalities, history of chest pain, CVD, marked HTN	Maximal treadmill or Two-step Double Master's	≥0.1 mV of ST depression 80 ms from the J point or exercise induced arrhythmia
Hollenberg	377 army officers	Known CHD	Maximal	≥1 mm ST
et al., 1985 (38)	Mean age 37		treadmill – USAFSAM Protocol	depression during or after exercise
	% male NR			or
				treadmill exercise score < 5 units
Boyle et al., 1987 (14)	1,174 employees from 2 factories in the U.K.	Symptoms of angina, orthopedic	Treadmill	Maximal ST/HR slope value of >13
	Mean age: NR Age range: 19-64	hypertension with retinopathy.		min 10 ⁻³
	95% male	fainting and afib		······································

* CABG, Coronary artery bypass graph surgery; CAD, Coronary artery disease; CHD, Coronary heart disease; CP, chest pain; CVD, Cardiovascular disease; DM, Diabetes mellitus; ECG, Electrocardiogram; ETT, Exercise tolerance test (treadmill); HTN, hypertension; J, Joules; mV, Millivolt; NR, Not reported; SD, Standard Deviation; VAMC, Veterans' Administration Medical Center; VD, vessel disease

	Asymptomatic	Frevalent Coro	nary near Di	sease (continueu)	
Preval ence of Abnor mal ETT	Definition of Abnormal Cardiac Catheterizatio n	Abnormal Catheteri- zations/ Total Catheteri- zations* (%)	Abnormal Catheteri- zations/ Abnormal ETT* (%)	Abnormal ETT and Abnormal Catheterizations / Total Screened Population* (%)	Quality Grading
22/349 6 (0.6%)	NR	10/15 (66.7%)	10/22 (45.5%)	10/3,496 (0.3%)	Fair
27/771 (3.5%)	NR	4/19 (21%)	4/27 (14.8%)	4/771 (0.5%) all cases were mild to moderate disease	Fair
45/377 (12%)	≥50% narrowing of the luminal diameter of major epicardial artery	1/10 (10%)	1/45 (2%)	1/377 (0.3%) 1 had 1-VD	Fair
3/377 (0.7%)			1/3		
68/1,17 4 (5.8%)	≥75% stenosis of epicardial artery	9/24 (37.5%)	9/68 (13.2%)	9/1,174 (0.8%) 1 had CABG	Fair

Author and Year Okin et al.,1988 (31)	Study Population 606 men in the Army Reserve at moderate to high risk by Framingham Risk score Mean age: NR Age: >40 years 100% male	Exclusion Criterion Known or suspected CHD or angina	Test Technique Modified Balke- Ware with radionuclide scintigram for an abnormal exercise ECG	Definition of Abnormal Exercise ECG ≥1 mm ST depression
Koistinen 1990 (15;75)	136 diabetics in Finland Mean age: 48 62% male	Clinical evidence of CHD, lipid lowering agents, DM less than 5 y, retinopathy and renal failure	Maximal bicycle ergometry and thallium scintigram	≥1 mm horizontal or downsloping ST depression
Dunn et al., 1991 (30)	1,930 patients referred to Cleveland Clinic Foundation for screening ETT in 1987-88(5.6% had history of chest pain) Mean age: 49 85% male	Known CAD	Symptom- limited exercise ECG, then thallium scintogram if exercise ECG abnormal	≥1 mm of horizontal or downsloping S ⁻ depression or arrhythmia

			-		
Prevalen ce of Abnorm al ETT	Definition of Abnormal Cardiac Catheterizatio n	Abnormal Catheteri- zations/ Total Catheteri- zations* (%)	Abnormal Catheteri- zations/ Abnormal ETT* (%)	Abnormal ETT and Abnormal Catheterizations / Total Screened Population* (%)	Quality Rating
10/606 (positive - abnormal exercise ECG and scintigram) (inconclu- sive- abnormal exercise ECG and normal scintigram	≥50% narrowing of the luminal diameter	7/10 (70%)	7/10 (70%)	7/606 (1.2%) 2 had 3-VD, 2 had 2-VD, 3 had 1-VD	Good
52/606					
40/136 (29%)	Significant ≥50% narrowing of the luminal diameter	12/34 (35%)	12/40 (30%)	12/136 (9%) 5 had 1VD 5-had 2 VD 2-had 3 VD	Fair
155/1,93 0 (8%)	≥50% blockage of any major vessel	CAD 25/41 (61%)	25/155 (16.1%)	25/1,930 (1.3%) 6 had CABG	Fair

Author and Year Massie et al., 1993 (18)	Study Population 226 men from the San Francisco VAMC – all had hypertension and at least 1 other cardiovascular risk factor Mean age (SD): 61 (8) 100% male	Exclusion Criterion Known cardiac disease history or symptoms, rest ECG abnormalities, paced rhythm, noncardiac limitation to exercise	Test Technique Standard Bruce with thallium scintigraphy	Definition of Abnormal Exercise ECG ≥0.1 mV of additional horizontal or downsloping S segment depression at 80 ms after the J point.
Davies et al., 1996 (23) Cameron et al., 1997 (25)	5,000 men from the United Kingdom Mean age: NR 100% male 229 Australians responding to questionnaire about chest pain Mean age: NR	NR Known CAD or negative screening questionnaire	Modified Balke Modified Bruce	1 mV of horizontal or downsloping depression persisting for ≥! complexes Flat ST segment depression ≥0.15 mV
Pilote et al., 1998 (28)	43% male 4,334 patients referred to Cleveland Clinic Foundation for screening ETT 19901993 Median age: 51 89% male	History of chest pain, heart failure, valvular or congenital heart disease, arrhythmia or digitalis use	Bruce or modified Bruce	≥1 mm horizontal or downsloping S ⁻ depression, ≥1 mm ST elevation in leads other than aVR or V1 drop in BP ≥ 10 mmHg, typical CP, failure to reach target heart rate

Prevale nce of Abnor mal ETT	Definition of Abnormal Cardiac Catheterizatio n	Abnormal Catheteri- zations/ Total Catheteri- zations* (%)	Abnormal Catheteri- zations/ Abnormal ETT* (%)	Abnormal ETT and Abnormal Catheterizations / Total Screened Population* (%)	
Abnorm al exercise ECG 67/226 (30%)	Intraluminal lesion of ≥50% diameter of vessel in 2 projections	Abnormal exercise ECG 14/26 (54%)	Abnormal exercise 14/67 (21%)	20/226 (9%) 6 had left main disease or 3-VD; 5 had 2-VD; 7 had 1-VD	Fair
Abnorm al scintigra m 41/226 (18%)	-	Abnormal scintigram 18/21 (86%)	Abnormal scintigram 18/29 (62%)	-	
162/5,0 00 (3.2%)	≥75% stenosis epicardial artery	67/86 (78%)	67/162 (41.4%)	67/5,000 (1.3%) 26 had CABG	Fair
Males 15/98 (15.3%) Female s 17/131 (13%)	NR	10/13 (77%)	10/32 (31%)	10/229 (4%) 4 had CABG	Fair
633/4,3 34 (15%)	CAD ≥1 coronary segment with ≥50% stenosis	71/126 (56%)	71/633 (11%)	71/4,334 (1.6%)	Poor

Table 2.Studies of the Use of Exercise Electrocardiogram to Detect
Asymptomatic Prevalent Coronary Heart Disease (continued)

Severe CAD -	19/126	19/633	19/4,334	Fair
left main	(15%)	(3%)	(0.4%)	
disease with				
≥50% stenosis				
or 3 vessel				
disease with				
≥70% stenosis				
or proximal				
LAD and 2V				
with ≥70%				
 stenosis				

Author and Year	Study Population	Exclusion Criterion	Test Technique	Definition of Abnormal Exercise ECG
Livschitz et al., 2000 (22)	4,900 male soldiers in the Israeli army ≥39 years of age	Angina, heart failure, valvular disease, congenital heart disease, or	Bruce	≥1 mV of horizontal or downsloping S ⁻ depression or ≥1.5 mV
	43(3) 100% male	arrhythmia		upsloping ST depression
Blumenthal et al., 2003 (17)	734 primarily white healthy siblings of individuals diagnosed with CAD before age 60 in Baltimore	Known CAD, limitations that precluded testing	Modified Bruce and thallium scintigraphy	NR for ETT
	Mean age: NR Age: < 60 years "Primorily molo"			
<u> </u>				

Prevalen ce of Abnorma I ETT	Definition of Abnormal Cardiac Catheterization	Abnormal Catheteri- zations/ Total Catheteri- zations* (%)	Abnormal Catheteri- zations/ Abnormal ETT* (%)	Abnormal ETT and Abnormal Catheterizations/ Total Screened Population* (%)
299/4,900 (6.1%)	NR	3/4 (75%)	3/299 (1%)	3/4900 (0.06%)
				1 had CABG
				2 had 1-VD
153/734 (21%)	Significant CAD intraluminal lesion of ≥50% diameter	41/105 (39%)	41/153 (27%)	41/734 (5.5%)
exercise ECG, scan, or both)	Hemodynamically significant CAD intraluminal lesion of ≥70% diameter	24/105 (23%)	24/153 (16%)	24/734 (3.3%)

*Percentages were calculated by the authors of this report.

			Mean Years	<u></u>		- <u>-</u> .
Author	Study		of Follow		Abnorma	al Test
Year	Population	Exclusion	-up	Technique	Definition	erevalenc
Giagn oni et al., 1983 (36)	514 factory workers in Italy Age range: 18- 65 73% male	Positive history and PE for CVD, rest BP ≥160/95, abnormal rest ECG	6	Submaximal supine cycle ergometry	≥1 mm of horizontal/ downsloping ST depress during or after exercise	NR
MRFI T Trial Resea rch Group , 1985(8) Rauta harju et al., 1986 (50)	6,205 MRFIT (multi-center cohort study – men in the upper 10% to 15% Framingham risk score distribution) Age range: 35-57 100% male	Clinical heart disease, life- limiting conditions, DBP ≥115, cholesterol ≥350	7	Submaximal	Computer code-ST seg depression 16 µV-s or more in leads CS5, aVL, aVF, V5 during or after exercise (in ECG with less than 6µV-s depress at rest)	12.2%
Gordo n et al., 1986 (41) Ekelu nd et al., 1989 (26)	3,640 white men in Lipid Research Clinics Prevalence Survey in US and Canada Mean age: 47 Age range: 35-59 100% male	Evidence of CHD based on history, rest ECG, and physician exam. Secondary HLP, BMI >32.1, BP ≥165/105. on anti-HTN or CV med, DM	8.1	Submaximal Modified Bruce	≥1 mm of ST depress or elevation/ computer- ST integral decreased or increased ≥10 µV-s from rest value	8.3%

* Events are CHD events unless otherwise indicated.

[†] CHD death.

[‡] All cause death

[§]AS, Aortic stenosis; aVL, Name of ECG lead; BMI, Body mass index; BP, Blood pressure; CAD, Coronary artery disease; CHD, Coronary heart disease; Chol, Cholesterol; CP, Chest pain; CV, Cardiovascular; CVD, Cardiovascular disease; DBP, Diastolic blood pressure; DM, Diabetes mellitus; ECG, Electrocardiogram; ETT, Exercise tolerance test); FBG, fasting blood glucose; HDL, High-density lipoprotein cholesterol; HR, Heart rate; HTN, Hypertension; LBBB, Left bundle branch block; LDL, Low-density lipoprotein cholesterol; MC, Minnesota Code; MRFIT,

Multiple Risk Factor Intervention Trial Research Group; NR, Not reported; PVC, Premature ventricular comples; RBBB, Right bundle branch block; SBP, Systolic block pressure; SVT, Supraventricular tachycardia; VO2, oxygen consumption; VT, Ventricular tachycardia

дәуі	mptomatic mutulu	ais (continueu)		
Cumulative Event Rate*	Adjusted Relative Risk for CHD Events with Abnormal ST Segment Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables
Normal ETT 3.4% Abnormal ETT 15.6%†	5.5 (2.8-11.2)	62	15	Age, SBP, smoking, coronary risk index
Normal ETT 2/1,000 person years† Abnormal ETT 7.6/1,000 person years†	3.5 (<i>P</i> <0.05) † 1.61 (<i>P</i> <0.01)‡	Not reported	36	Age, DBP, cholesterol, number of cigarettes smoked daily

<i>Placebo group</i> Normal ETT 13/1,000	<i>Placebo group</i> 5.7 (2.7-12.2) †	30	7.1	Age, LDL, HDL, SBP, smoking, family history
person years†	33			
Abnormal ETT 1.9/1,000 person years†	(1.8-5.9) ‡			
Cholestyramine group	Cholestyramine group 4.9			
Normal ETT 7.2/1,000	(2.2-10.8)†			
person years†	2.9 (1.6-5.2) ‡			
Abnormal ETT 1.5/1,000 person years†				

			Mean Years	<u></u>	Abnorm	nal Test
Author and Year	Study Population	Exclusion	of Follow- up	Technique	Definition	Prevalenc
Fleg et al., 1990 (19)	407 residents of Baltimore, Maryland (mainly white) Mean age	NR	4.6	Maximal treadmill with Thallium Modified Balke	≥1 mm of horizontal/ downsloping during or after exercise	Abnormal ECG only 16.0%
	(3D): 00 (11) Age range: 40- 90					Abnormal Thallium only 14%
	71% male					Both tests abnormal 6.0%
Okin et al., 1991 (40)	3,168 participants in the Framingham Offspring Study Mean age (SD): 44 (10) Age range: 17-70 48% male	Medical contraindicati ons to exercise, history of myocardial infarction, CHF, valvular disease, syncope, conduction abnormalities, digoxin use, atrial fibrillation	4.3	Standard Bruce	ST segment corrected for heart rate index >1.6 µV/beats/ min or abnormal rate recovery loop	416/3168 13% (either tes abnormal)
Siscovick et al., 1991 (12)	3,617 white men in the Lipid Research Clinics Prevalence Survey Mean age: NR Age range: 35-59 100% male	Clinical evidence of CHD or CHF on history, various rest ECG abnormalities	7.4	Modified Bruce Submaximal	Visual code - ≥1 mm ST depression or elevation or Computer code - ≥10 µV/sec	6.6%

	Adjusted			
	Relative Risk for CHD Events with Abnormal		Positive Predictive Value of	Relative Risk
Cumulative Event Rate*	ST Segment Response	Sensitivity for CHD Events	Abnormal ST Response	Adjusted for the Following Variables
Both normal 7%	1			Age, sex, HTN, FBG, total cholesterol, BML smoking
Abnormal ECG only 12%	(P < 0.05)	40	24	exercise duration
,.	1.4 (NS)	N/A	N/A	
Abnormal Thallium	()			
only 3%	3.6 (1.6-8.1)	28	48	
Both tests abnormal 48%				
Both normal 1.6%	1			Age, sex, smoking, DBP, total
Either test abnormal	1.6 (1.1-2.5)	23%	4%	cholesterol, FBG, LVH on ECG
4.1% Both tests abnormal 9.8%	2.7 (1.8-4.0)	8%	10%	
Overall 2% (for CHD events occurring during	2.6 (1.3 – 5.2) (for CHD events occurring during exercise)	18%	5%	Age, LDL, HDL, smoking, physical activity, workload achieved, family history of CHD, BMI,

alcohol consumption

Association between Abnormal ST Segment Response to Exercise and Coronary Heart Disease Events in Asymptomatic Individuals (continued) Table 3.

during

exercise)

		<u> </u>	Mean Years			
Author	Study		of Follow-		Abnorn	
and Year	Population	Exclusion	up	Technique	Definition	Prevalence
Blumenth al et al., 1996 (32)	264 healthy siblings of individuals with	Known CAD, corti- costeriods,	6.2	Bruce	≥1 mV (≥2 mm for women) of	normal
	CAD before age 60 in Baltimore	collagen vascular disease,		Thallium scintigraphy	horizontal or downsloping depression	+exercise ECG 5.4%
	Mean age (SD): 46 (8) Age range: 37-	decreased life expectancy			in 3 consecutive beats during	+scan 18.1%
	59	functional status			exercise or first 3 min of	+exercise ECG and
	69% male	limitations			recovery	SUAIT 4.070
Okin et al., 1996 (39)	5,940 men in the usual care group of MRFIT	No evidence of CHD by history, physical of	7	Submaximal treadmill	ST segment corrected for heart rate index	729/5,940 (12.3%)
	Mean age: NR Age range: 35- 57	rest ECG			>1.6µV/ beats/min	
	100% male					
Katzel et al., 1999 (29)	170 healthy sedentary obese men living in the Baltimore- Washington, DC area (96% white) Mean age: NR Age range: 45-79	history or laboratory evidence of CAD, DM, HTN, hyperlipidemia	7.3	Maximal Bruce	≥1 mm of horizontal or downsloping ST segment depression in 2 or more leads	37/170 (22%)
	100% male					

Asy	mptomatic mulv	iduais (continued	1)	
Cumulative Event Rate*	Adjusted Relative Risk for CHD Events with Abnormal ST Segment Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables
Normal 3%	1			Age, sex
+exercise ECG 7%	1.5 (0.2-12.5)	N/A	N/A	-
+scan 13%	3.6	63%	20%	
	(1.1-11.4)			
+exercise ECG and scan 50%	14.5 (4.2-50.2)	32%	50%	-
	2.6	26%	E0/	Ass DDD shall
	(2.4-5.4)†	30 76	3%	smoking
5.4%†				
Overall 18%	4.23 (2.03-8.83)	55%	46%	age, BMI, maximal VO2, fasting glucose

			Mean Years of		Abnorm	al Test
Author and Year	Study Population	Exclusion	Follow- up	Technique	Definition	Prevalenc e
Gibbons et al., 2000 (33)	25,927 patients of a preventive medicine clinic in Texas (mainly white) Mean age: 42.9 Age range 20-82 100% male	Evident CHD, severe AS, acute systemic illness, uncontrolled atrial or ventricular arrhythmias, pericarditis, myocarditis, thrombophlebitis or exercise limiting orthopedic problems	8.4	Maximal treadmill Modified Blake	CP and ≥ 1mm ST segment depression or elevation also exercise induced- dec ≥10mm in SBP, SBP >250, DBP >1 20, VT, LBBB, RBBB, SVT	No risk factors, 3.0% >1 risk factor, 7.1%

ASy	inpromatic mutur	duais (continued)	
Cumulative Event Rate	Adjusted Relative Risk for CHD Events with Abnormal ST Segment Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables
No risk factor Normal ETT 0.08/1000 person years†	21 (6.9-63.3)†	60	2.2	Age
Abnormal ETT 2.8/1000 person years†				
>1 risk factor Normal ETT 0.5/1000 person years†	9†	61	7.7	·
Abnormal ETT 7.6/1000 person years†				

			Me Ye	an ars of			Abnorr	nal Test
Author and Year	Study Population	n Exclusion	Fol	-wo	Techniq	ue	Definition	Prevalence
Josephson et al., 1990 (11)	1,083 participants in the Baltimore	History of angina or heart failure, Q	7.9	Modif	ied Balke	Nor	mal	
Rywik et al., 2002 (21)	Longitudinal Study of Aging Mean age (SD): 52(18) 57% male	wave on rest ECG, valvular disease, use of anti- arrhythmic drugs, those who did not achieve 85% of max heart rate				MC mm dep with dow ST com any	11.1-≥1 J point ression flat or vnsloping segment in st nplexes in lead	20%
						MC dep rest wor MC exe	11.5 ST ression at that sens to 11.1 during rcise	5.5%
						MC Hor dow ST of 0	11.2- izontal or /nsloping depression .5-1.0 mm	7%
						MC poir dep ≥1 r Ups	11.4-J nt ression of nm with loping ST	11.5%

Exe Asy	rcise and Corona mptomatic Individ	ry Heart Disease duals (continued)	Events in)	
Cumulative Event Rate*	Adjusted Relative Risk for CHD Events with Abnormal ST Segment Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables
Male 4% Female 3%	1	Male 74%	Male 16%	Age, cholesterol,
		Female 68%	Female 7%	duration
Male 17% Female 8%	OR 2.7 (1.6-4.7)			
Male 17% Female 11%	OR 2.7 (1.05-7.10)			
Male 10% Female 5%	OR 1.8 (0.6-5.4)			
Male 17% Female 3%	OR 1.3 (0.6-2.9)			

Table 3 Association between Abnormal ST Segment Response to

Author and	Ctudy		Mean Years of			
Year	Population	Exclusion	-wollor- up	Technique	Abnormal Te	st
Jouven and Ducimetiere, 2000 (45)	6,101 Frenchmen Paris Civil Service Age range: 42-53 100% male	in Known or suspected CVD, SBP≥180 at rest, rest ECG abnormality	23	Bicycle ergometry	J-point depression of at least 1 mm with a flat or downsloping ST segment during exercise or recovery	4.4%
Laukkanen et al., 2001 (20)	1,769 study participants, population in Kupio Ischemic Heart Disease Study base sample of Finnish men Mean age (SD): 52 (5.2) 100% male	Known CHD or symptoms suggestive of CHD	10	Maximal Bicycle ergometry	>1 mm ST depression during exercise	10.7 %
Rutter et al., 2002 (13)	86 diabetics in the U.K. Mean age (SD): 62(7) Age range: 46- 74 72% male	History of CAD	2.8	treadmill	> 1 mm of horizontal or downsloping ST segment depression for 3 consecutive beats	(52 %)
Mora et al., 2003(42)	2994 women enrolled in the Lipid Research Clinics Prevalence Study Age range 30-80 0% male	Pregnancy or significant cardiovascula r disease	20.3	Maximal Bruce protocol	≥1mm horizontal or downsloping ST depression at 0.08 seconds after the J point during recovery or exercise	4.7 %

Asymptomatic Individuals (continued)							
Cumulative Event Rate*	Adjusted Relative Risk for CHD Events with Abnormal ST Segment Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables			
Normal ETT 6.4% Abnormal ETT 16.7%†	2.6 (1.93-3.59) †	10	17-25	Age, BMI, HR at rest, smoking, physical activity, DM, total chol, PVC			
Normal ETT 9.2% 2.4%† Abnormal 15.3% 7.9%†	1.7 (1.1-2.6) 3.5 (1.9-6.5) †	16	15	Age, examination year, smoking, SBP, alcohol consumption, BMI, max oxygen uptake, DM, LDL, HDL			
Both normal and abnormal ETT 17%	21 (2-204)	100%	20%	Ankle brachial index, microalbuminuria, Framingham 10-y CHD risk >30%, fibrinogen			
Both normal and abnormal ETT 5%† 14%‡	0.88 (0.48-1.61) † 0.69 (0.45-1.04) ‡	-	-	Age, smoking, diabetes,family history of premature heart disease, obesity, HDL, LDL, triglycerides, hypertension			

- <i></i>	Study	Exclusion	Mean Years of	Test	Definition of	Prevalence
Ekelund et	3.106 (healthy	Men with CVD	8.5	Modified Bruce	HR during stage 2	Increase of 2
al., 1988 (35)	group of white men) in Lipid Research	sx and/or HTN were analyzed separately		submaximal	of ETT and exercise time	SD in stage 2 HR
	Clinics Prevalence Survey in US and Canada					Decrease of 2 SD in time on the treadmill
	Age range: 30- 69					
	<u>10</u> 0% male					
Lauer et al., 1996 (44)	1,575 subjects in Framingham Offspring Study (predominantly white)	Prevalent CAD, inability to reach stage 2 in Bruce protocol, use of β -blockers at time of ETT	7.7	Bruce protocol submaximal	Failure to achieve age- and sex- predicted target heart rate on ETT	21%
	wean age: 43					
	100% male			_		

Table 4. Association between Exercise Predictors and Coronary Heart

Disease Events in Asymptomatic Individuals[§]

* Events are CHD events unless otherwise indicated.

† CHD death.

[‡] All cause death [§] BMI, Body mass index; CAD, Coronary artery disease; CHD, Coronary heart disease; Chol, Cholesterol; COPD, Chronic obstructive pulmonary disease; CVD, Cardiovascular disease; DM, Diabetes mellitus; ECG, Electrocardiogram; ESRD, End-stage renal disease; ETT, Exercise tolerance test; HDL, High-density lipoprotein cholesterol; HR, Heart rate; HTN, Hypertension; LDL, Low-density lipoprotein cholesterol; METS, unit of metabolic work; PVC, Premature ventricular complex; PVD, Premature ventricular depolarizations (synonym to PVC); SBP, Systolic blood pressure; SD, Standard Deviation; SES, Socio-economic status; VEA, Ventricular ectopic arrhythmia; VFib, Ventricular fibrillation; VPC, Ventricular premature contractions; VT, Ventricular tachycardia

			Positive	Relative Risk
Cumulative	Relative Risk for CHD	Sensitivity for CHD	Predictive Value of Abnormal ST	Adjusted for the Following
Event Rate	Events with Positive Test	Events	Response	Variables
0.26-1.69%†	3.2 (1.5-6.7)	NR	NR	Age, smoking, HDL, LDL, SBP
	2.8 (1.3-6.1)			
3% for those who reached target heart rate (All cause death)	No significant association of predictor to all cause death 1.75 (1.11-2.74)†	46%	14%	Age, ST segment response, physical activity, BMI, smoking, HTN, HTN medication, DM
6% for those who failed to reach heart rate‡				total cholesterol/HDL

Table 4.Association between Exercise Predictors and Coronary Heart
Disease Events in Asymptomatic Individuals (continued)

			Mean Years of			
Author Year	Study Population	Exclusion Criteria	Follow- up	Test Technique	Definition of Abnormal Test	Prevalence of Predictor
Wei et al., 1999 (48) Blair et al	25,714 patients at a preventive med clinic in	History of cancer, BMI < 18.5, age < 20 and	24	Maximal treadmill	Low fitness using age-based METS cut points on ETT	Normal weight: 10%
1996 (49)	Texas Aerobics Center	those with < 1 year of follow- up			UTLIT	Overweight: 24%
	Longitudinal Study (>95% white), 10% of men with known CVD					Obese: 50%
	Mean age: 43.8					
	100% male					
Cole et al., 2000 (34)	5, 234 in Lipid Research Clinics Prevalence Survey in US and Canada	Age < 30, use of β -blockers, digoxin, antiarrhythmic agents or nitrates,	12	Bruce or modified Bruce submaximal	Abnormal HR recovery defined as heart rate change of 42 beats/min or less from peak	33%
	Mean age: >30 years	history of cardiovascular disease,			exercise to that measured 2 min later	
	39% male	unable to reach stage 2				
Jouven and Ducimetier 2000 (45)	6, 101 Frenchmen in Paris civil service	Known or suspected CVD, SBP ≥180 at rest, rest ECG	23	Bicycle ergometry	PVCs constituting more than 10% of all ventricular depolarizations	2.3%
	Age range: 42- 53	abnormality			during exercise	
	100% male					

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Table 4.Association between Exercise Predictors and Coronary HeartDisease Events in Asymptomatic Individuals (continued)

Cumulative Event Rate	Relative R Events with Segment	isk for CHD Abnormal ST Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables
Overall 1.7/1,000 person	1.7 (1.1-2.5)†	1.6 (1.3-2.1)‡	36%	4.6%	DM, cholesterol, HTN, current smoking, history
years†	1.9 (1.4-2.5)†	1.7 (1.4-2.0)‡	52%	5.4%	of CVD, abnormal ECG at rest, age, BMI parental
	2.0 (1.5-3.4)†	2.3 (1.5-3.4)‡	79%	3.4%	history of CVD, examination year

Table 4.Association between Exercise Predictors and Coronary HeartDisease Events in Asymptomatic Individuals (continued)

Normal heart rate recovery 4% death Abnormal heart rate recovery 10% death	1.95 (1.11-3.42)† 1.55 (1.22-1.98)‡	54%	10%	Age, sex, BMI, ethnicity, SBP, HTN medication, exercise habits, physical fitness, smoking, DM, lipids, ST segment response, HR, chronotropic index, SES
Normal ETT 6.4% Abnormal ETT 16.1%†	2.53 (1.65-3.88)†	5%†	17%†	Age, BMI, heart rate, SBP, tobacco, level of physical activity, DM, total cholesterol, presence or absence of PVD before or after exercise

Author	Study	Exclusion	Mean Ye of Folic	ears ow- Test	Definition of	Preva-lence
Year	Population	criterion	up	Technique	Abnormal Test	of Predictor
Morshedi- Meibodi et al., 2002 (47)	2,967 participants in Framingham Off spring Study	Prevalent CVD, COPD, use of digoxin or β- blockers, rest ECG	15	Bruce submaximal	Heart rate recovery index – decline in peak heart rate to time 2 minute of <42	N/A
	Mean age (SD): 43 (10)	abnormali- ties, inability to complete			beats per minute	
	47% male	stage 1 of exercise				
Rywik et al., 2002 (21)	1,083 participants in the Baltimore Longitudinal Study of Aging Mean age (SD): 52(18) 57% male	History of angina or HF, Q wave on rest ECG, valvular disease, use of antiarrhythmic drugs, those who did not achieve 85% of max heart rate	7.9	Modified Balke	Duration of exercise	NA
Frolkis et al., 2003 (46)	29,244 persons referred to Cleveland Clinic for ETT Mean age (SD): 56 (11) 70% male	Age <30, symptomatic heart failure, use of digoxin, valvular disease, ESRD, pacer, atrial fibrillation, heart block, freq VEA at rest, heart transplant, concurrent evaluation for an arrhythmia	5.3	Bruce protocol submaximal	Frequent VEA - ≥7VPC's/ minute, ventricular bigeminyor trigeminy, ventricular couplets or triplets, VT, ventricular flutter, torsade de pointes or VFib	No VEA Freq VEA during recovery 2% Freq VEA during exercise 3%

Table 4.Association between Exercise Predictors and Coronary Heart
Disease Events in Asymptomatic Individuals (continued)

			Positive	Relative Risk
	Relative Risk for CHD	Sensitivity	Predictive Value	Adjusted for
Cumulative	Events with Abnormal ST	for CHD	of Abnormal ST	the Following
Event Rate	Segment Response	Events	Response	Variables
Overall 7.2%	1.1	NA	NA	Age, BMI,
	(0.8-1.5)			smoking,
				SBP, DBP,
	0.8			anti-HTN
	(0.5-1.1) ‡			medication,
				DM, total
				cholesterol,
				HDL, resting
				heart rate
				and peak
	0.07		NID	
Overall 7 %		INK	INE	Aye, cholesterol
	(0.79-0.90) (Ear CHD event for 1 minute			Cholesterol,
	increase in evercise			seament
	duration)			changes
	ddration)			Ghàngeo
5%‡	1			Age, sex, DM,
				HTN, smoking,
11%‡	Freq VEA during recovery 1.5	3%	12%	prior CAD,
	(1.1-1.9)‡			medication use,
			- • •	BMI, resting
9%‡	Freq VEA during exercise 1.1	4%	9%	heart rate SBP,
	(0.9-1.3)‡			Si segment
				changes,
				chronotropic
				incompetence,
				abnormal neart
				rate recovery,
				peak exercise
. <u> </u>			· · ••••••••	capacity

Table 4.Association between Exercise Predictors and Coronary HeartDisease Events in Asymptomatic Individuals (continued)

Author Year	Study Population	Exclusion criterion	Mean Years of Follow- up	Test Technique	Definition of Abnormal Test	Prevalence of Predictor
Mora et al., 2003	2994 women enrolled in the Lipid Research Clinics Prevalence Study	Pregnancy or significant cardiovascular disease	20.3	Maximal Bruce protocol	Low exercise capacity (< 7.5 METS) and low heart rate recovery (<55 beats/minute)	31%
	Age range 30-60					
	0% male					
Gulati et al., 2003	5721 women from the Chicago area (86% white) Mean age 52 years 0% male	Self reported CHD, Percutaneous coronary intervention, coronary bypass surgery, congestive heart failure	9	Maximum Bruce protocol	Exercise capacity in METS	-

Table 4.Association between Exercise Predictors and Coronary Heart
Disease Events in Asymptomatic Individuals (continued)

Cumulati ve Event Rate	Relative Risk for CHD Events with Abnormal ST Segment Response	Sensitivity for CHD Events	Positive Predictive Value of Abnormal ST Response	Relative Risk Adjusted for the Following Variables
Both	3.52†	71%	11%	Age, smoking,
normal	(1.57-7.86)			diabetes,family
and				history of
abnormal	2.11‡			premature heart
ETT	(1.47-3.04)			disease, obesity,
5%†				HDL, LDL,
14%‡				triglycerides,
				hypertension
3.2%‡	0.83	-	-	Framingham
	(0.78-0.89)			Risk Score
	for each 1 MET increase in			
	exercise capacity			

Table 4.Association between Exercise Predictors and Coronary Heart
Disease Events in Asymptomatic Individuals (continued)