

TASK FORCES

Task Force I: Determination of Prognosis in Patients With Ischemic Heart Disease

DAVID B. PRYOR, MD, FACC, CHAIRMAN, ROBERT A. BRUCE, MD, FACC, BERNARD R. CHAITMAN, MD, FACC, LLOYD FISHER, PhD, FACC, JERZY GAJEWSKI, MD, PhD, KARL E. HAMMERMEISTER, MD, FACC, STEPHEN G. PAUKER, MD, FACC, JOSEPH STOKES III, MD

I. Introduction

This report summarizes information regarding morbidity and mortality in ischemic heart disease as it relates to insurability and employability. *Insurability* requires the assessment and evaluation of risks that may relate to life, disability and health. *Employability* should consider three aspects: 1) the capability of the individual to perform the job, 2) the risk to the individual of performing the job, and 3) the risk to society if the individual is performing the job.

This report focuses primarily on the assessment of risk of patients with chronic ischemic heart disease, including definite or probable angina pectoris, previous myocardial infarction, cardiac arrest or documented significant coronary artery obstruction. It focuses primarily on chronic disease outcomes and does not address the early prognosis of patients presenting with acute myocardial infarction. In addition, the risk to society posed by ischemic heart disease events occurring on the job is considered for individuals without manifestations of coronary artery disease. Prognostic risk focuses primarily on survival, but also encompasses subsequent myocardial infarction. The effects of specific therapies on prognosis and changes in the natural history of the disease are also considered.

II. Sources of Information

Three types of studies that have recently been extensively reviewed (1) concern prognosis in patients with ischemic heart disease: 1) U.S. population statistics, 2) cohort or observational studies, and 3) randomized clinical trials.

U.S. population statistics. These data, obtained from national surveys, census figures and publications from the American Heart Association (2-5), provide limited information for the evaluation of specific individuals, and there are hazards in extrapolating results from specific studies to national statistics (6-8).

Cohort or observation studies. These studies define the population through community or regional surveillance. Findings from these studies can be extrapolated to the U.S. population because the study population is generally repre-

sentative of the U.S. population as a whole. The findings from several large studies (the Framingham Study, Tecumseh Study, Chicago Peoples Gas Company Study, Minneapolis Professional and Business Men, U.S. Railroad Workers studies and the Chicago Western Electric Company study) largely agree with each other. The risk of developing coronary heart disease among asymptomatic individuals as it relates to the initial assessment has been defined by these studies. Estimates from the Framingham Study form the basis of the Coronary Risk Handbook (9).

Most of the detailed information regarding the assessment of risk of individual patients with manifest ischemic heart disease is derived from specific cohort studies of individuals with the disease. In general, these studies provide the most detailed assessment of the importance of characteristics influencing prognosis but suffer from the selection bias introduced from the enrollment process and criterion selection. The Coronary Artery Surgery Study (CASS) trial included a large collaborative patient registry of all individuals catheterized at participating centers (10). Collaborative registries of patients with acute myocardial infarction include the Myocardial Infarction Limitation of Size (MILIS) Study (11), the Multicenter Postinfarction Trial (MPIT) (12) and those coordinated at San Diego (13) and Stanford (14); however, this report focuses primarily on nonacute disease.

Voluntary collaborative registries include the National Heart, Lung, and Blood Institute-Percutaneous Transluminal Coronary Angioplasty (PTCA) registry of patients undergoing percutaneous transluminal coronary angioplasty (15). There are large regionally-based databases describing complete assessment and outcome of patients undergoing cardiac catheterization or other specific procedures at one or more institutions. These include the Duke Database for Cardiovascular Disease (16), the Seattle Heart Watch Study (17), the Cleveland Clinic Registry (18), the University of Alabama Registry (19) and the Emory University Registry (20). Prognostic studies of specific groups of patients undergoing tests or procedures include those from Cedars-Sinai Hospital (21).

Randomized controlled trials. These trials have evaluated not only the outcome of coronary bypass surgery and medical

Table 1. Prognostic Characteristics From the Initial Assessment

| Characteristics | Analysis | |
|-----------------------------------|--------------------|--------------|
| | Univariate | Multivariate |
| Age | CA, CL, D, I, S | CA, D, S |
| Gender | CA, D, I | CA, D |
| Risk factors | | |
| Smoking | CA, CL, D, I | CA |
| Hypercholesterolemia | D, I | |
| Hypertension | CA, CL, D, I, S, V | CA, CL, V |
| Diabetes | CL, D, S | CL |
| Symptoms | | |
| Pain type | D, S | D |
| Duration | CL, D, S | CL |
| Course | CA, CL, D, V | CA, D, V |
| Nocturnal | D | |
| Frequency | D, S | S |
| Myocardial damage | | |
| Congestive heart failure or class | CA, CL, D, S | CL, D, S |
| ST-T wave changes | CL, D, S, V | CL, D, S, V |
| Ventricular gallop | D, S | D |
| Cardiomegaly | CA, CL, D, S | CA, D, S |
| Premature ventricular complexes | D, S | D |
| History of myocardial infarction | CA, D, V | D, V |
| Q waves on electrocardiogram | CL, D, S | CL, D |
| Digitalis use | CA, S | |
| Diuretic use | CA, S | |
| Rales | D, S | |
| Heart murmur | D, S | |
| Left ventricular hypertrophy | CL, D, S | CL, S |
| Conduction abnormalities | CL, D, S | CL, D |
| Peripheral vascular disease | CL, D, E, S | CL, D |

CA = Coronary Artery Surgery Study Trial and Registry (10,94-98); CL = Cleveland Clinic Registry (18,103); D = Duke University Database (16,99-101); E = European Coronary Surgery Study (90-93); I = Life Insurance Studies (1,104); S = Seattle Heart Center Watch Registry (17,22,102); V = Veterans Administration Cooperative Trial (87-89).

therapy but also changes in the natural history of ischemic heart disease. Assessments of prognosis provided by these studies are limited by selection bias introduced by who is enrolled in the trial and difficulties in generalizing beyond these subjects of patients. This report summarizes the three largest trials in patients with stable angina: the Veterans Administration Cooperative Trial, the European Coronary Surgery Study and the Coronary Artery Surgery Study.

III. Characterization of Prognosis

A. Prognostic Characteristics From the Initial Assessment

Table 1 shows significant prognostic variables from the initial assessment that have been found to be important in at least two of the seven major studies. Where the relation has been examined for the characteristic alone and found to be significant, an appropriate notation has been made in the

univariate column. Where multivariate analyses have been performed including at least several of the characteristics (uncommon for the randomized clinical trials) and the characteristics have remained important, an appropriate notation has been made in the multivariate column. The prognostic evaluation of a patient with coronary artery disease begins with the physician's initial assessment. Decisions to perform noninvasive testing or cardiac catheterization should be considered in light of what these tests add to the physician's initial assessment.

Clinical perspective versus life insurance perspective on prognosis. Many different characteristics from the physician's initial assessment can be used to predict survival. Their importance as they relate to prognosis depends in part on how an individual's prognosis is considered. The prognostic risk from a life insurance perspective compares the risk of an individual with coronary artery disease with that which would be expected for an individual of similar age and gender. Clinicians generally consider the prognosis from the perspective of the importance of individual characteristics among patients with established disease. This difference in perspective regarding prognosis influences the importance assigned to characteristics from the initial assessment.

Factors affecting prognosis. Among patients with established coronary artery disease, clinical prognosis worsens with age. However, when considered from a life insurance perspective, the relation reverses (relative mortality in ischemic heart disease patients is considerably higher in younger than in older age groups) because the assigned risk of the individual with coronary artery disease is related to the expected risk in a comparable healthy population. Consequently, age is an important prognostic characteristic and of fundamental importance when considered from the life insurance perspective.

Other important prognostic characteristics from the initial assessment include the patient's gender, risk factors, symptoms, evidence of myocardial damage and evidence of associated vascular disease. The importance of gender is marginal in patients with established disease, although this characteristic is very important in operative mortality with coronary artery bypass grafting, most likely related to the smaller size of coronary vessels in women. Risk factors are particularly important in relation to the risk of development of coronary artery disease, but their contribution to prognosis is less important among patients with established disease (16,22). When examined by themselves, they clearly identify subgroups at higher risk, but when considered in conjunction with other characteristics, their contribution is less important. Similarly, psychological constructs such as the type A personality appear to be important in the development of the disease, but evidence of their importance as independent prognostic variables is not convincing.

The presence of significant myocardial damage has been found in all studies to be an important predictor of outcome.

Table 2. Prognostic Characteristics: Treadmill Exercise Testing

| Characteristic | Studies (reference no.) | | |
|-------------------------|---------------------------|-----------------------|--------------------------------|
| | Univariate Analysis | Multivariate Analysis | Add to Cardiac Catheterization |
| ST segment change | 23, 24, 27, 105-111 | 23, 94, 105 | 105 |
| Treadmill angina | 23, 24, 105-107 | 105 | 105 |
| Treadmill time or stage | 23, 94, 105, 107, 109-112 | 94, 105, 107, 113 | 94, 105, 113 |
| Maximal heart rate | 23, 94, 105, 111, 114 | 23, 94 | 94 |
| Ventricular arrhythmias | 23, 105, 115, 116 | 23 | — |
| Anginal heart rate | 117 | — | — |
| Exertional hypotension | 24, 118 | — | — |

In essence, this characteristic reflects the presence and extent of previous myocardial infarction and is often associated with symptoms of congestive heart failure.

Anginal symptoms identify individuals with myocardium at risk, and their tempo reflects the severity of the risk. Longstanding angina reflects longstanding coronary artery disease and, because the disease is progressive in nature, suggests a worsening anatomic severity.

B. Prognostic Characteristics Elicited by Noninvasive Testing

Noninvasive testing has been able to identify high and low risk subgroups of patients with chronic coronary artery disease. In general, studies are relatively small in size and, to provide sufficient events, have often combined end points to include not only death but also myocardial infarction and coronary artery bypass grafting.

The treadmill exercise test. This test has been the most carefully studied and excellent reviews are available (23-26). The studies cited in Table 2 indicate that among the many exercise variables evaluated, the most important prognostic indicators are maximal exercise capacity, increases in systolic blood pressure, and ST segment displacement (depression and elevation) (27). Although uncommon, exertional hypotension and ventricular tachycardia also identify a high risk subgroup of patients. The magnitude of the additional

prognostic information provided by exercise testing is considerable in relation to the initial assessment and comparatively modest where catheterization has been performed (28,29).

Nuclear studies. Important prognostic characteristics obtained with radionuclide angiography and with thallium scintigraphy are shown in Tables 3 and 4, respectively. The radionuclide exercise ejection fraction appears to be the most important prognostic variable. In patients with a normal rest ejection fraction, the change in ejection fraction with exercise and the peak exercise ejection fraction provide virtually identical prognostic information. Perfusion defects in the distribution of two or more coronary arteries, and perhaps the quantitative estimate of the magnitude of hypoperfusion or the uptake of thallium in the lung appear to be the most important thallium prognostic variables. The contributions of radionuclide angiography and thallium scintigraphy to prognostic assessment are similar.

Ambulatory electrocardiography. This has also been performed in large series of patients with acute myocardial infarction and chronic angina. In patients with acute myocardial infarction, two large observational studies (11,12) have shown that frequent premature ventricular complexes on a predischARGE 24 h ambulatory ECG are predictive of impaired prognosis independently of left ventricular function. Similar observations have been made in patients with

Table 3. Prognostic Characteristics: Radionuclide Angiography

| Characteristic | Studies (reference no.) | |
|-----------------------------|-------------------------|-----------------------|
| | Univariate Analysis | Multivariate Analysis |
| Exercise ejection fraction | 119, 120 | 120 |
| Change in ejection fraction | 121 | |
| Rest ejection fraction | 119-121 | 119 |
| Angina | | |
| ECG changes | | |
| Wall motion abnormalities | 120 | |
| Exercise time | 120 | |
| Rest end-diastolic volume | 119 | |

Table 4. Prognostic Characteristics: Thallium Scintigraphy

| Characteristic | Studies (reference no.) | |
|---------------------------------|-------------------------|-----------------------|
| | Univariate Analysis | Multivariate Analysis |
| Transient defects | 21, 122-124 | 21, 122, 124 |
| Total or fixed defects | 21, 122, 125 | 123 |
| Degree of hypoperfusion | 21 | 21 |
| Lung thallium uptake | 126 | 126 |
| Exercise ST changes | 124, 126 | 124, 126 |
| Heart rate changes | 124 | 124 |
| Exercise ventricular arrhythmia | 124 | |

Table 5. Prognostic Characteristics: Cardiac Catheterization

| | Studies | |
|--|-----------------------|-----------------------|
| | Univariate Analysis | Multivariate Analysis |
| Ventricular function | | |
| Ejection fraction | CA, D, I, S | CA, D, S |
| LV score or wall motion abnormalities | CA, CL, D, S | CA, CL, D |
| Anatomy | | |
| Left main disease | CA, D, I, V | CA, D, V |
| Number of diseased vessels | CA, CL, D, E, I, S, V | CA, CL, D, S |
| Proximal LAD | CA, D, S | CA, D |
| Other proximal disease or jeopardy score | CA, D | CA, D |
| Mitral regurgitation | CA, D, S | CA, D, S |
| LVEDP | CL, D, S | CL, D, S |

CA = CASS Trial and Registry (10,94-98); CL = Cleveland Clinic Registry (18,103); D = Duke University Database (16,99-101); E = European Coronary Surgery Study (90-93); I = Life Insurance studies (1, 104); LAD = Left anterior descending coronary artery; LV = Left ventricular coronary artery; LVEDP = Left ventricular end-diastolic pressure; S = Seattle Heart Center Watch Registry (17, 22, 102); V = VA Trial (87-89).

congestive heart failure (30), but this relation has not been found in patients with angina requiring cardiac catheterization (31). Premature ventricular complexes occur more frequently in patients with reduced left ventricular function, and their prognostic information overlaps with measurements of left ventricular function (31). Evidence of silent ischemia has also been shown to be related to prognosis, although the data are not fully developed.

Exercise echocardiography. This study may also identify patients with a poor prognosis through assessment of wall motion changes, however, the data are not fully developed.

C. Prognostic Characteristics From the Cardiac Catheterization

Prognostically important characteristics from cardiac catheterization are shown in Table 5. Information from the physician's initial assessment and cardiac catheterization is shown in Table 6.

The prognostically most important characteristics from the catheterization reflect left ventricular function and coronary anatomy. Left ventricular function is usually expressed as the ejection fraction. Anatomic characteristics reflecting myocardial jeopardy include the degree of left main stenosis, the number of significantly diseased major coronary vessels and the presence of proximal disease in the major coronary arteries, particularly of the left anterior descending coronary artery. The severity of mitral regurgitation and elevated left ventricular end-diastolic pressure are also prognostically important.

Important characteristics from the initial assessment combined with the cardiac catheterization are shown in Table 6. The most important predictors of survival are left ventricular function and coronary anatomy. Patients with unstable angina or poor left ventricular function and severe congestive heart

failure have an especially unfavorable prognosis. The presence of other vascular disease, mitral regurgitation and age have also been found to be significant predictors of outcome when considered with information from cardiac catheterization. When examined by themselves, risk factors clearly identify subgroups at higher risk but when considered in conjunction with the characteristics shown in Table 6, their contribution is less important. The importance of gender is marginal in patients with established disease.

Prognostic estimates are most discriminating when all characteristics are considered. For example, consider the prognosis of two patients with three vessel disease and a normal ejection fraction. One is 64, has frequent angina, resting ST depression, peripheral vascular disease, previous myocardial infarction, an ejection fraction of 51% and a 95%

Table 6. Summary of Important Prognostic Characteristics: Initial Assessment and Catheterization

| | Studies | |
|--------------------------|---------------------|-----------------------|
| | Univariate Analysis | Multivariate Analysis |
| Age | CA, CL, D, I, S | CA, CL, D, S |
| Gender | CA, D, I | CA, D |
| Risk factors | CA, CL, D, I, S, V | CA, CL, V |
| Symptoms | CA, CL, D, S, V | CA, CL, D, S, V |
| Myocardial damage | All | All |
| Vascular disease | CA, CL, D, E, S | CA, CL, D, E |
| Conduction abnormalities | CL, D, S | CL, D, S |
| Anatomy | All | All |
| LV function | CA, CL, D, I, S, V | CA, CL, D, S, V |
| Mitral regurgitation | CA, D, S | D, S |

CA = Coronary Artery Surgery Study Trial and Registry (10,94-98); CL = Cleveland Clinic Registry (18,103); D = Duke University Database (16, 99-101); E = European Coronary Surgery Study (90-93); I = Life Insurance studies (1,104); LV = left ventricular; S = Seattle Heart Center Watch Registry (17,22,102); V = Veterans Administration Trial (87-89).

proximal left anterior descending coronary artery lesion. The other is 51, has infrequent angina, no peripheral vascular disease or previous myocardial infarction, an ejection fraction of 64% and a distal 75% left anterior descending coronary artery lesion. Expected survival on medical therapy is quite different. Five year estimates from the Duke Database (32) are 42% for the first patient and 92% for the second.

IV. Effects of Therapy on Prognosis

Studies of prognosis in ischemic heart disease must consider the ameliorating effects of therapy, including medical therapy, coronary artery bypass grafting, coronary angioplasty and changes over time in the natural history of the disease.

A. Medical Therapy

The influence of risk factor modification and drug therapy on the prognosis of patients with established disease has been reviewed (1,33). In general, serum cholesterol, hypertension and particularly smoking behavior worsen the prognosis of patients with manifest ischemic heart disease. However, the influence of these characteristics on *recurrent* cardiac events is modest compared with their influence on the incidence of *initial* coronary events (34-41). Although therapy directed at modifying these risk factors remains a prudent recommendation, definitive demonstration of substantially improved survival with risk factor modification in patients with chronic stable angina is limited (40).

Trials of drug therapy in modifying medical prognosis have been largely limited to patients with acute myocardial infarction or unstable angina. Evidence for an improved outcome, due to beta-blocker therapy, is more convincing after acute myocardial infarction (42-44) than in chronic angina (26,45,46). Two studies (47,48) have shown dramatic benefit of aspirin therapy in patients hospitalized with unstable angina. Some benefit for calcium channel blocker therapy may also be present (49). Vasodilator therapy may also improve survival in patients with congestive heart failure including some patients with coronary artery disease (50,51). The evidence for the use of antiarrhythmic therapy is controversial.

B. Coronary Artery Bypass Graft Surgery

The value of surgery in modifying prognosis has been excellently reviewed (1,52-54). In general, the magnitude of the improvement in prognosis is greatest in patients with anatomically extensive disease (left main disease and three vessel disease) or high risk based on clinical evaluation. Improvement is sustained in the first 7 to 8 years after surgery, diminishing thereafter as grafts become occluded (55,56,57). The influence of repeat bypass grafting on prog-

nosis is unknown at present. The improved patency of internal mammary artery grafting compared with saphenous vein bypass procedures may improve or extend survival benefits with surgery (58,59).

Primary characteristics influencing operative mortality include age, body size (identified by gender in many studies), hypertension, prior coronary artery bypass grafting, history of smoking, left ventricular function, extent of anatomical disease, the presence of unstable angina or recent myocardial infarction, cerebral or peripheral vascular disease, and renal or pulmonary dysfunction (60-62) (Hammermeister KE. Seattle Heart Watch 1988 [personal communication]. Pryor DB. Duke Database 1988 [personal communication]). Long-term outcome after coronary artery bypass grafting is influenced by the same characteristics that define the risk in medically-treated patients except that coronary anatomy is much less important.

C. Coronary Angioplasty

The value of angioplasty in modifying prognosis in patients with coronary artery disease is unknown (63,64). The major limitation with respect to long-term prognosis appears to be restenosis occurring within 6 months in approximately one-third of treated patients (65,66).

D. Changes in the Natural History of Ischemic Heart Disease

There is overwhelming evidence that the natural history of coronary artery disease in the United States is improving over time. A recent comprehensive review describes a 42% decline in the age-adjusted mortality rates in the U.S. population between 1963 and 1985 (67). Outcomes have improved for patients with acute myocardial infarction (34,68) and those undergoing coronary artery bypass surgery (69). Improvements for patients with chronic stable angina treated medically are more modest (34,69,70).

V. Consideration of Occupations Posing a Risk to Society

Clinicians are frequently asked to "certify" that it is appropriate for an individual with ischemic heart disease to return to work. Such decisions are especially important in public safety officers, firefighters, pilots, critical process operators (e.g., nuclear power plant operators) and professional drivers. A significant cardiac event such as cardiac arrest or myocardial infarction occurring at a "critical" time in such occupations may adversely affect not only the individual but also the public or coworkers. In "certifying" whether it is appropriate for such individuals to work, the physician should consider not only the risk to the individual, but also the risk to society.

Prognostic concerns about coronary heart disease have led to the mandatory retirement of individuals with demonstrated disease, as well as individuals without manifest disease. These two groups will be considered separately with respect to both the risk to the individual and the risk to society.

A. Asymptomatic Individuals Without Manifest Coronary Disease

In general, there is no convincing evidence that employment in a specific occupation worsens an individual's risk for the occurrence of a cardiac event. A possible exception is firefighting, which entails bursts of heavy physical activity and exposure to heat and psychological stress. The risk of cardiac events in individuals without manifest coronary disease can be estimated with use of standard risk factor equations such as the Coronary Risk Handbook (9) based on the Framingham Study.

Epidemiologic studies suggest that habitual physical exercise may offer some protection against primary or secondary events of coronary heart disease and associated mortality but "falls short of proving" (71) this. The evidence also suggests that the effect of exercise training is outweighed by the other risk factors. There has also been some demonstration that emotional stress and sudden bursts of activity may increase the risk of sudden death, particularly in sedentary individuals (72,73).

Risk factor profiling using techniques such as the Coronary Risk Handbook are accurate and well substantiated. Screening programs to detect individuals without manifest coronary disease should consider such strategies rather than relying solely on individual characteristics such as age, gender, smoking, and others. Exercise testing can further improve the identification of individuals with one or more risk factors (74-77) and can be applied economically to specific occupations (78). In some cases, such as for airline pilots, it may be appropriate to consider further evaluation with radionuclide procedures or coronary angiography, or both, for individuals at increased risk (79-84).

B. Symptomatic Individuals

Whether it is appropriate for symptomatic individuals to return to occupations posing a risk to the public is often difficult to determine. From a prognostic standpoint, the goal is to determine whether the job is likely to place the individual under a stress that would otherwise be unlikely to occur. However, it is difficult to duplicate or simulate on-the-job conditions, such as those encountered in firefighting (85,86), in a standard ECG stress laboratory.

The decision should also consider the status of the patient's disease. The risk of death or infarction is substantially higher in individuals with manifest or symptomatic coronary disease than in those without manifest disease. The

Table 7. Prognostic Characteristics Classified by Pathophysiologic Construct and Weighted According to Power in Predicting Prognosis for Patients With Manifest Coronary Artery Disease

| Pathophysiologic Construct | Weight |
|--|--------|
| Myocardial function | |
| Congestive heart failure | ++++ |
| Functional class | +++ |
| Diuretic use | ++ |
| Digitalis use | ++ |
| Cardiomegaly | +++ |
| S ₃ gallop | +++ |
| Prior myocardial infarction | ++ |
| Exercise systolic pressure | +++ |
| Exercise capacity | ++++ |
| Ventricular arrhythmias | ++ |
| Ejection fraction | ++++ |
| Wall motion score or assessment | ++++ |
| Left ventricular end-diastolic pressure | ++ |
| Myocardial jeopardy | |
| Angina frequency | + |
| Duration of angina | + |
| Level of exercise inducing ischemia | +++ |
| Degree of exercise induced ST segment depression | +++ |
| Exercise systolic pressure | +++ |
| Presence and extent of reversible thallium uptake | +++ |
| Number and distribution of coronary vessels with obstruction | +++ |
| Myocardial ischemia | |
| Rest or nocturnal angina | +++ |
| Progressive angina | ++ |
| Unstable angina | +++ |
| Failure of angina to respond to nitroglycerin | ++ |
| ST segment depression on rest ECG | +++ |

prognostic value of silent ischemia, defined as ST segment depression without chest pain, in patients without manifest coronary artery disease is unknown. For certain groups of patients, such as those with unstable angina, return to work is clearly inappropriate in a setting where the occurrence of a significant cardiac event would place the public at risk. In contrast, individuals without clinical or angiographic evidence of myocardial ischemia after coronary artery bypass grafting have an excellent prognosis (82).

VI. Conclusions

1. Variables from the clinical assessment, noninvasive tests and cardiac catheterization that are important for estimating prognosis include descriptors of myocardial function, myocardial jeopardy and myocardial ischemia (Table 7).

2. Prognosis can be estimated from the clinical examination (history, physical examination, ECG, chest roentgenogram) combined when indicated with one or more noninvasive studies (e.g., exercise test, radionuclide angiography, myocardial imaging or echocardiography) and does not usually require cardiac catheterization except when dictated by regulations or recommendations.

3. Most occupations do not increase the risk of coronary events. In patients with manifest coronary disease, continued employment should be permitted when patients are functionally able to perform the job.

4. Prognosis should be estimated for occupations in which sudden disability might endanger others. For patients with manifest disease, this evaluation should occur at yearly intervals, or more frequently if required by regulation or recommendation. The cost effectiveness of these approaches needs to be evaluated.

5. Prognosis should also be estimated for patients in occupations requiring sudden or sustained high level physical effort or exposure to extremes of hot or cold or to hypoxia, hypercarbia, carbon monoxide or stimuli producing sudden bursts of autonomic activity.

6. Prognosis need not be estimated for individuals without manifest coronary disease, except perhaps for those with occupations in which sudden disability might endanger others. In the latter group, noninvasive assessment may be of value. In some instances, cardiac catheterization may be required to satisfy regulations or recommendations.

7. Although the advisability of return to full-time work of individuals with manifest coronary disease can usually be established a priori, a trial work period may sometimes be necessary.

References

- Singer RB, Gajewski J. Cardiovascular Diseases I. In: Lew EA, Gajewski J, eds. *Medical Risks—Trends in Mortality*. New York: Praeger, 1989:6-1-59.
- Advance report of final mortality statistics, 1984. NCHS Monthly Vital Statistics Report 1986;35:1-44.
- Thornberry OT, Wilson RW, Golden P. Health promotion and disease prevention provisional data from the National Health Interview Survey: United States, January—June 1985. NCHS Advancedata 1986;119:1-16.
- American Heart Association. 1987 Heart Facts. Dallas, TX: American Heart Association, 1987:1-31.
- American Heart Association. 1986 Heart Facts. Dallas, TX: American Heart Association, 1986:1-31.
- Califf RM, Lee KL, Harrell FE Jr, Kimm SY, Gruffman S, Rosati RA. Assessment of the use of the age- and sex-specific United States population as a control group for analysis of survival in coronary artery disease. *Am J Cardiol* 1982;50:1279-82.
- Lawrie GM, Morris GC Jr, Howell JF, Tredici TD, Chapman DW. Improved survival after 5 years in 1,444 patients after coronary bypass surgery. *Am J Cardiol* 1978;42:709-15.
- Hurst JW, King SB, Logue RB. Value of coronary bypass surgery. *Am J Cardiol* 1978;42:308-29.
- American Heart Association. *Coronary Risk Handbook: Estimating Risk of Coronary Heart Disease in Daily Practice*. New York: American Heart Association, 1973:1-35.
- Mock MB, Fisher L, Killip T, et al. The natural history of nonoperated patients with ischemic heart disease: the CASS experience. In: Hammermeister KE, ed. *Coronary Bypass Surgery: the Late Results*. New York: Praeger, 1983:83-98.
- Mukharji J, Rude RE, Poole WK. Risk factors for sudden death after acute myocardial infarction: two-year follow-up. *Am J Cardiol* 1984;54:31-6.
- Bigger JT, Fleiss JL, Miller JP, Rolnitzky LM, Multicenter Post-Infarction Research Group. The relationships among ventricular arrhythmias, left ventricular dysfunction, and mortality in the 2 years after myocardial infarction. *Circulation* 1984;69:250-8.
- Madsen EB, Gilpin EA, Ahnve S, Henning H, Ross J Jr. Prediction of functional capacity and use of exercise testing for predicting risk after acute infarction. *Am J Cardiol* 1985;56:839-45.
- DeBusk RF, Kraemer HC, Nash E, Berger WE III, Lew H. Stepwise risk stratification soon after acute myocardial infarction. *Am J Cardiol* 1983;52:1161-6.
- Kent KM, Bentivoglio LG, Block PB, et al. Long-term efficacy of percutaneous transluminal coronary angioplasty (PTCA): report from the National Heart, Lung, and Blood Institute PTCA Registry. *Am J Cardiol* 1984;53:27C-31C.
- Harris PJ, Harrell FE, Lee KL, Behar VS, Rosati RA. Survival in medically treated coronary artery disease. *Circulation* 1979;60:1259-69.
- DeRouen TA, Hammermeister KE, Dodge HT. Comparisons of the effects on survival after coronary artery surgery in subgroups of patients from the Seattle Heart Watch. *Circulation* 1981;63:537-45.
- Proudfit WL, Bruschke AVG, MacMillan JP, Williams GW, Sones FM Jr. Fifteen-year survival study of patients with obstructive coronary artery disease. *Circulation* 1983;68:986-97.
- Kouchoukos NT, Oberman A, Kirklin JW, et al. Coronary bypass surgery: analysis of factors affecting hospital mortality. *Circulation* 1980;62(suppl I):I-84-I-9.
- Jones EL, Craver JM, King SB III. Clinical, anatomic and functional descriptors influencing morbidity, survival and adequacy of revascularization. *Ann Surg* 1980;192:390-402.
- Ladenheim ML, Kotler TS, Pollock BH, Berman DS, Diamond GA. Incremental prognostic power of clinical history, exercise electrocardiography and myocardial perfusion scintigraphy in suspected coronary artery disease. *Am J Cardiol* 1987;59:270-7.
- Hammermeister KE, DeRouen TA, Zia M, Dodge HT. Survival of medically treated coronary artery disease patients in the Seattle Heart Watch angiography registry. In: Hammermeister KE, ed. *Coronary Bypass Surgery: the Late Results*. New York: Praeger, 1983:167-94.
- Peduzzi P, Hultgren H, Thomsen J, Angell W. Prognostic value of baseline exercise tests. *Prog Cardiovasc Dis* 1986;28:285-92.
- Ellestad MH, Allen WH, Stuart RJ. Diagnostic and prognostic information derived from exercise testing. *Cardiovasc Clin* 1978;9:33-55.
- Chaitman BR. The changing role of the exercise electrocardiogram as a diagnostic and prognostic test for chronic ischemic heart disease. *J Am Coll Cardiol* 1986;8:1195-210.
- Froelicher VF. *Exercise and the Heart: Clinical Concepts*. Chicago: Year Book Medical Publishers, 1987:157-64.
- Bruce RA, Fisher LD, Pettinger M, Weiner DA, Chaitman BR. ST segment elevation with exercise: a marker for poor ventricular function and poor prognosis. *Circulation* 1988;77:897-905.
- Harrell FE Jr, Califf RM, Pryor DB, Lee KL, Rosati RA. Evaluating the yield of medical tests. *JAMA* 1982;247:2543-6.
- Mark DB, Hlatky MA, Harrell FE Jr, Lee KL, Califf RM, Pryor DB. Exercise treadmill score for predicting prognosis in coronary artery disease. *Ann Intern Med* 1987;106:793-800.
- Cleland JFG, Dargie HJ, Ford I. Mortality in heart failure: clinical variables of prognostic value. *Br Heart J* 1987;58:572-82.
- Califf RM, McKinnis RA, Burks J. Prognostic implications of ventricular arrhythmias during 24 hour ambulatory monitoring in patients undergoing catheterization for coronary artery disease. *Am J Cardiol* 1982;50:23-31.
- Califf RM, Pryor DB, Greenfield JC Jr. Beyond randomized clinical trials: applying clinical experience in the treatment of patients with coronary artery disease. *Circulation* 1986;74:1191-4.

33. Goldman GJ, Pichard AD. The natural history of coronary artery disease: does medical therapy improve the prognosis? *Prog Cardiovasc Dis* 1983;25:513-52.
34. Elveback LR, Connolly DC. Coronary heart disease in residents of Rochester, Minnesota. V. Prognosis of patients with coronary heart disease based on initial manifestation. *Mayo Clin Proc* 1985;60:305-11.
35. Coronary Drug Project Research Group. Natural history of myocardial infarction in the Coronary Drug Project: long-term prognostic importance of serum lipid levels. *Am J Cardiol* 1978;42:489-98.
36. Leren P. The effect of plasma cholesterol lowering diet in male survivors of myocardial infarction. *Acta Med Scand* 1966;466(suppl):1-90.
37. Little JA, Shanoff HM, Roe RD, et al. Studies of male survivors of myocardial infarction. IV. Serum lipids and five-year survival. *Circulation* 1965;31:854-62.
38. Weinblatt E, Shapiro S, Frank CW, et al. Prognosis of men after first myocardial infarction: mortality and first recurrence in relation to selected parameters. *Am J Public Health* 1968;58:1329-47.
39. Frank CW, Weinblatt E, Shapiro S. Angina pectoris in men: prognostic significance of selected medical factors. *Circulation* 1973;47:509-17.
40. Kallio V, Hamalainen H, Hakkila J, et al. Reduction in sudden deaths by a multifactorial intervention programme after acute myocardial infarction. *Lancet* 1979;2:1091-4.
41. Shaw PJ, Bates D, Carlidge NEF, et al. Neurological complications of coronary artery bypass graft surgery: six month follow-up study. *Br Med J* 1986;293:165-7.
42. Norwegian Multicenter Study Group. Timolol-induced reduction in mortality and reinfarction in patients surviving acute myocardial infarction. *N Engl J Med* 1981;304:801-7.
43. Hjalmarson A, Herlitz J, Malek I, et al. Effect on mortality of metoprolol in acute myocardial infarction. A double-blind randomized trial. *Lancet* 1981;2:823-7.
44. Beta-Blocker Heart Attack Trial Research Group. A randomized trial of propranolol in patients with myocardial infarction. I. Mortality results. *JAMA* 1982;247:1707-14.
45. Lambert DMD. Beta blockers and life expectancy in ischemic heart disease. *Lancet* 1972;1:793.
46. Lambert DMD. Effect of propranolol on mortality in patients with angina. *Postgrad Med J* 1976;52(suppl IV):57-60.
47. Lewis HD, Davis JW, Archibald DG, et al. Protective effects of aspirin against acute myocardial infarction and death in men with unstable angina. *N Engl J Med* 1983;309:396-403.
48. Cairns JA, Gent M, Singer J, et al. Aspirin, Sulfapyrazone, or both in unstable angina. *N Engl J Med* 1985;313:1369-75.
49. Gibson RS, Young PM, Boden WM, Schechtman K, Roberts R, Diltiazem Reinfarction Study Group. Prognostic significance and beneficial effect of Diltiazem on the incidence of early recurrent ischemia after non-Q-wave myocardial infarction: results from the multicenter Diltiazem Reinfarction Study. *Am J Cardiol* 1987;60:203-9.
50. Cohn JN, Archibald DG, Ziesche S, et al. Effect of vasodilator therapy on mortality in chronic congestive heart failure. *N Engl J Med* 1986;314:1547-52.
51. Consensus Trial Study Group. Effects of Enalapril on mortality in severe congestive heart failure. *N Engl J Med* 1987;316:1429-73.
52. Frye RL, Fisher L, Schaff HV, Gersh BJ, Vlietstra RE, Mock MD. Randomized trials in coronary artery bypass surgery. *Prog Cardiovasc Dis* 1978;30:1-22.
53. Gersh BJ, Califf RM, Loop FD, Akins CW, Pryor DB, Takaro TC. Coronary bypass surgery in chronic stable angina. *Circulation* 1989;79(suppl 1):146-59.
54. Hammermeister KE. The effect of coronary bypass surgery on survival. *Prog Cardiovasc Dis* 1983;25:297-334.
55. Califf RM, Harrell FE Jr, Lee KL, et al. Changing efficacy of coronary revascularization: implications for patient selection. *Circulation* 1988;78(suppl 1):1-185-1-9.
56. Campeau L, Enjalbert M, Lesperance J, Vaislic C, Grondin CM, Bourassa MG. Atherosclerosis and late closure of aortocoronary saphenous vein grafts: sequential angiographic studies at 2 weeks, 1 year, 5 to 7 years, and 10 to 12 years after surgery. *Circulation* 1983;68(suppl II):II-1-II-7.
57. Bourassa MG, Fisher LD, Campeau L, Gillespie MJ, McConney M, Lesperance J. Long-term fate of bypass grafts: the coronary artery surgery study (CASS) and Montreal Heart Institute Experiences. *Circulation* 1985;72(suppl V):V-71-V-8.
58. Rankin JS, Newman GE, Bashore TM, et al. Clinical and angiographic assessment of complex mammary artery bypass grafting. *J Thorac Cardiovasc Surg* 1986;92:832-46.
59. Loop FD, Lytle BW, Cosgrove DM, et al. Influence of the internal mammary artery on 10-year survival and other cardiac events. *N Engl J Med* 1986;314:1-6.
60. Fisher LD, Kennedy JW, Davis KB, et al. Association of sex, physical size, and operative mortality after coronary artery bypass in the Coronary Artery Surgery Study (CASS). *J Thorac Cardiovasc Surg* 1982;84:334-41.
61. Kennedy JW, Kaiser GC, Fisher LD, et al. Clinical and angiographic predictors of operative mortality from the Collaborative Study in Coronary Artery Surgery (CASS). *Circulation* 1981;63:793-801.
62. Kennedy JW, Kaiser GC, Fisher LD, et al. Multivariate discriminant analysis of the clinical and angiographic predictors of operative mortality from the Collaborative Study in Coronary Artery Surgery (CASS). *J Thorac Cardiovasc Surg* 1980;80:876-87.
63. Gruentzig AR, King SB III, Schlumpf M, Siegenthaler W. Long-term follow-up after percutaneous transluminal coronary angioplasty: the early Zurich experience. *N Engl J Med* 1987;316:1127-32.
64. Vlietstra RE, Holmes DR. PTCA: Percutaneous Transluminal Coronary Angioplasty. Philadelphia: FA Davis, 1987:17.
65. Blackshear JL, O'Callaghan WG, Califf RM. Medical approaches to prevention of restenosis after coronary angioplasty. *J Am Coll Cardiol* 1987;9:834-47.
66. Schwartz L, Bourassa MG, Lesperance J, et al. Aspirin and dipyridamole in the prevention of restenosis after percutaneous transluminal coronary angioplasty. *N Engl J Med* 1988;318:1714-9.
67. Thom JT, Maurer J. Time trends for coronary heart disease mortality and morbidity. In: *The Influence of Medical Care*. Higgins MW, Luepker RF, eds. New York: Oxford University Press, 1988:7-15.
68. Goldberg RJ, Gore JM, Alpert JS, Dalen JE. Incidence and case fatality rates of acute myocardial infarction 1975-1984: The Worcester Heart Attack Study. *Am Heart J* 1988;115:761-7.
69. Pryor DB, Harrell FE Jr, Rankin JS, et al. The changing survival benefits of coronary revascularization over time. *Circulation* 1987;76(suppl V):V-13-V-21.
70. Pryor DB, Harrell FE Jr, Lee KL, Califf RM, Rosati RA. An improving prognosis over time in medically treated patients with coronary artery disease. *Am J Cardiol* 1983;52:444-8.
71. Leon AS. Physical activity levels and coronary heart disease. *Med Clin North Am* 1985;69:3-20.
72. Eliot RS, Buell JC. Role of emotions and stress in the genesis of sudden death. *J Am Coll Cardiol* 1985;5:95B-8B.
73. Siscovick DS, Weiss NS, Fletcher RH, Lasky T. The incidence of primary cardiac arrest during vigorous exercise. *N Engl J Med* 1984;311:878-81.
74. Bruce RA, Fisher LD. Exercise-enhanced risk factors for coronary heart disease vs. age as criteria for mandatory retirement of healthy pilots. *Aviat Space Environ Med* 1987;58:792-8.
75. Bruce RA, Hossack KF, DeRouen TA, Hofer V. Enhanced risk assessment for primary coronary heart disease events by maximal exercise testings: 10 years' experience of Seattle Heart Watch. *J Am Coll Cardiol* 1983;2:565-73.
76. Uhl GS, Froelicher V. Screening for asymptomatic coronary artery disease. *J Am Coll Cardiol* 1983;3:946-55.

77. Bruce RA, Fisher LD, Hossack KF. Validation of exercise-enhanced risk assessment of coronary heart disease events: longitudinal changes in incidence in Seattle community practice. *J Am Coll Cardiol* 1985; 5:875-81.
78. Hartley LH, Herd JA, Day WC, Abusamra J, Howes B. An exercise testing program for large populations. *JAMA* 1979;241:269-71.
79. Hopkirk JA, Uhl GS, Hickman JR, Fischer J, Medina A. Discriminant value of clinical and exercise variables in detecting significant coronary artery disease in asymptomatic men. *J Am Coll Cardiol* 1984;3:887-94.
80. Froelicher VF, Thompson AJ, Wolthuis R, et al. Angiographic findings in asymptomatic aircrewmembers with electrocardiographic abnormalities. *Am J Cardiol* 1977;39:32-8.
81. McHenry PL, O'Donnell J, Morris SN, Jordan JJ. The abnormal exercise electrocardiogram in apparently healthy men: a predictor of angina pectoris as an initial coronary event during long-term follow-up. *Circulation* 1984;70:547-51.
82. Chaitman BR, Davis KB, Dodge HT, et al. Should airline pilots be eligible to resume active flight status after coronary bypass surgery?: a CASS Registry study. *J Am Coll Cardiol* 1986;8:1318-24.
83. Booze CF Jr, Staggs CM. A comparison of postmortem coronary atherosclerosis findings in general aviation pilot fatalities. *Aviat Space Environ Med* 1987;58:297-300.
84. Ross J Jr, Pepine CJ, Brandenburg RO, et al. Guidelines for coronary angiography: a report of the American College of Cardiology/American Heart Association Task Force on Assessment of Diagnostic and Therapeutic Cardiovascular Procedures (Subcommittee on Coronary Angiography). *J Am Coll Cardiol* 1987;10:935-50.
85. Davis PO, Dotson CO. Heart rate responses to fire fighting activities. *Ambulator Electrocardiol* 1978;1:15-8.
86. Puterbaugh JS, Lawyer CH. Cardiovascular effects of an exercise program: a controlled study among firemen. *J Occup Med* 1983;25:581-6.
87. Hultgren H, Peduzzi P, Pierpont G. Symptom severity prognosis. *Prog Cardiovasc Dis* 1986;28:273-8.
88. Detre K, Hultgren H, Takaro T. Veterans Administration Cooperative Study of Surgery for Coronary Arterial Occlusive Disease. *Am J Cardiol* 1977;40:212-24.
89. Veterans Administration Coronary Artery Bypass Surgery Cooperative Study Group. Eleven-year survival in the Veterans Administration randomized trial of coronary bypass surgery for stable angina. *N Engl J Med* 1984;311:1333-9.
90. Varnauskas E, European Coronary Surgery Study Group. Survival, myocardial infarction, and employment status in a prospective randomized study of coronary bypass surgery. *Circulation* 1985;72(suppl V): V-90-101.
91. European Coronary Surgery Study Group. Coronary-artery bypass surgery in stable angina pectoris: survival at two years. *Lancet* 1979;889-93.
92. European Coronary Surgery Study Group. Long-term results of prospective randomized study of coronary artery bypass surgery in stable angina pectoris. *Lancet* 1982;1173-80.
93. Varnauskas E, Olsson SB. The European Multicenter CABG Trial. In: Yu PN, Goodwin JF, eds. *Progress in Cardiology*. Philadelphia, Chicago: Lea & Febiger, 1977:83-9.
94. Weiner DA, Ryan TJ, McCabe CH, et al. Prognostic importance of a clinical profile and exercise test in medically treated patients with coronary artery disease. *J Am Coll Cardiol* 1984;3:772-9.
95. Cameron A, Schwartz MJ, Kronmal RA, Kosinski AS. Prevalence and significance of atrial fibrillation in coronary artery disease (CASS Registry). *Am J Cardiol* 1988;61:714-7.
96. CASS Principal Investigators and Their Associates. Coronary Artery Surgery Study (CASS): a randomized trial of coronary artery bypass surgery: survival data. *Circulation* 1983;68:939-50.
97. CASS Principal Investigators and Their Associates. Myocardial infarction and mortality in the coronary artery surgery study (CASS) randomized trial. *N Engl J Med* 1984;310:750-8.
98. Passamani E, Davis KB, Gillespie MJ, Killip T, CASS Principal Investigators and Their Associates. A randomized trial of coronary artery bypass surgery: survival of patients with a low ejection fraction. *N Engl J Med* 1985;312:1665-71.
99. Harrell FE Jr, Lee KL, Califf RM, Pryor DB, Rosati RA. Regression modelling strategies for improved prognostic prediction. *Stat Med* 1984; 3:143-52.
100. Califf RM, Mark DB, Harrell FE, et al. Importance of clinical measures of ischemia in the prognosis of patients with documented coronary artery disease. *J Am Coll Cardiol* 1988;11:20-6.
101. Califf RM, Phillips HR III, Hindman MC, et al. Prognostic value of a coronary artery jeopardy score. *J Am Coll Cardiol* 1985;5:1055-63.
102. Hammermeister KE, DeRouen TA, Dodge HT. Variables predictive of survival in patients with coronary disease. Selection by univariate and multivariate analyses from the clinical electrocardiographic, exercise, arteriographic and quantitative angiographic evolution. *Circulation* 1979; 59:421-30.
103. Proudfit WL, Brusckge AVG, Sones FM Jr. Natural history of obstructive coronary artery disease: ten-year study of 601 nonsurgical cases. *Prog Cardiovasc Dis* 1978;21:53-78.
104. 1983 Medical Impairment Study. Chicago: Society of Actuaries and Association of Life Insurance Medical Directors of America 1986;1:64-5.
105. Mark DB, Hlatky MA, Harrell FE, Lee KL, Califf RM, Pryor DB. Exercise treadmill score for predicting prognosis in coronary artery disease. *Ann Intern Med* 1987;106:793-800.
106. Weiner DA, Ryan TJ, McCabe CH, et al. Significance of silent myocardial ischemia during exercise testing in patients with coronary artery disease. *Am J Cardiol* 1987;59:725-9.
107. Wyns W, Musschaert-Beauthier E, Domburg Rv, et al. Prognostic value of symptom limited exercise testing in men with a high prevalence of coronary artery disease. *Eur Heart J* 1985;6:939-45.
108. Ellestad MH, Wan MKC. Predictive implications of stress testing: follow-up of 2700 subjects after maximum treadmill stress testing. *Circulation* 1975;51:363-9.
109. McNeer JF, Margolis JR, Lee KL. The role of the exercise test in the evaluation of patients for ischemic heart disease. *Circulation* 1978; 57:64-70.
110. Podrid PJ, Graboys TB, Lown B. Prognosis of medically treated patients with coronary-artery disease with profound ST-segment depression during exercise testing. *N Engl J Med* 1981;305:1111-6.
111. Dagenais GR, Rouleau JR, Christen A, Fabia J. Survival of patients with a strongly positive exercise electrocardiogram. *Circulation* 1982;65:452-6.
112. Brusckge AVG, Proudfit WL, Sones FM. Progress study of 590 consecutive nonsurgical cases of coronary disease followed 5-9 years I. arteriographic correlations. *Circulation* 1973;47:1147-53.
113. Weiner DA, Ryan TJ, McCabe CH, et al. Value of exercise testing in determining the risk classification and the response to coronary artery bypass grafting in three-vessel coronary artery disease: a report from the Coronary Artery Surgery Study (CASS) registry. *Am J Cardiol* 1987; 60:262-6.
114. Gohlke H, Samek L, Betz P, Roskamm H. Exercise testing provides additional prognostic information in angiographically defined subgroups of patients with coronary artery disease. *Circulation* 1983;68:979-85.
115. Derry JM, Abouantoun S, Wyns W. Incidence and prognostic implications of severe ventricular arrhythmias during maximal exercise testing. *Cardiology* 1981;68:35-43.
116. Califf RM, McKinnis RA, McNeer JF, et al. Prognostic value of ventricular arrhythmias associated with treadmill exercise testing in patients studied with cardiac catheterization for suspected ischemic heart disease. *J Am Coll Cardiol* 1983;2:1060-7.
117. Hayet M, Kellerman JJ. The angina pectoris threshold heart rate as a prognostic sign. *Cardiology* 1981;68:78-83.
118. Bruce RA, DeRouen TA, Peterson DR, et al. Noninvasive predictors of

- sudden cardiac death in men with coronary heart disease. *Am J Cardiol* 1977;39:833-40.
119. Iskandrian A, Hakki A, Schwartz JS, Kay H, Mattleman S, Kane S. Prognostic implications of rest and exercise radionuclide ventriculography in patients with suspected or proven coronary heart disease. *Int J Cardiol* 1984;6:707-18.
120. Pryor DB, Harrell FE, Lee KL, et al. Prognostic indicators from radionuclide angiography in medically treated patients with coronary artery disease. *Am J Cardiol* 1984;53:18-22.
121. Borer JS, Wallis J, Hochreiter C, Holmes J, Moses JW. Prognostic value of left ventricular dysfunction at rest and during exercise in patients with coronary artery disease. *Adv Cardiol* 1986;34:179-85.
122. Brown KA, Boucher CA, Okada RD, et al. Prognostic value of exercise thallium-201 imaging in patients presenting for evaluation of chest pain. *J Am Coll Cardiol* 1983;1:994-1001.
123. Iskandrian AS, Hakki A, Kane-Marsch S. Exercise thallium-201 scintigraphy in men with nondiagnostic exercise electrocardiograms. *Arch Intern Med* 1986;146:2189-93.
124. Kaul S, Lilly DR, Gascho JA, et al. Prognostic utility of the exercise thallium-201 test in ambulatory patients with chest pain: comparison with cardiac catheterization. *Circulation* 1988;77:745-58.
125. Harris PJ, Lee KL, Harrell FE, Behar VS, Rosati RA. Outcome in medically treated coronary artery disease: ischemic events: nonfatal infarction and death. *Circulation* 1980;62:718-26.
126. Gill JB, Ruddy TD, Newell JB, Finkelstein DM, Strauss HW, Boucher CA. Prognostic importance of thallium uptake by the lungs during exercise in coronary artery disease. *N Engl J Med* 1987;317:1485-9.

Task Force II: Determination of Occupational Working Capacity in Patients With Ischemic Heart Disease

WILLIAM L. HASKELL, PHD, FACC, CHAIRMAN, NORMAN BRACHFELD, MD, FACC, ROBERT A. BRUCE, MD, FACC, PAUL O. DAVIS, PHD, CHARLES A. DENNIS, MD, FACC, SAMUEL M. FOX III, MD, FACC, PETER HANSON, MD, ARTHUR S. LEON, MD, FACC

Introduction

The physical, metabolic and cardiovascular demands of occupations in industrialized countries have substantially declined during the past century. Many jobs that once required substantial physical effort are now performed by machines, robots or computers. This decrease in job-related energy demand is especially apparent in occupations performed by people over age 40, who, given some seniority, rarely perform physical tasks exceeding a peak energy expenditure of 5 kcal/min or 3.5 METs, where 1 MET = energy expenditure sitting at rest (1). Nonetheless, the physical stress of employment is still the greatest challenge to the cardiovascular system of many patients with ischemic heart disease.

For many sedentary jobs posing limited psychological or environmental demands, the adequacy of the patient's physical working capacity can be assessed by a medical history, physical examination and symptom-limited exercise testing. However, as the physical demands of the job tasks increase to include exercise of widely varying intensities and types, or substantial psychological or environmental stress, assessment of physical working capacity becomes more complex.

In patients with ischemic heart disease, the focus of occupational work evaluation is to determine whether or not the increase in cardiac demands produced by physical, psychological and environmental stressors will exceed the threshold for a "safe working capacity." The challenge to

the physician is to obtain an accurate, valid and reliable determination of this capacity.

Factors Influencing Metabolic and Cardiovascular Demands During Work

Physical exertion increases metabolic and cardiac demands roughly in proportion to the absolute intensity of the exertion. The magnitude of effort required for a particular task by an individual is related to the individual's physical working capacity. Therefore, the metabolic and cardiac demands on an individual are related to characteristics of the task, such as the work intensity, type of work, size of muscle mass involved, work-rest cycle and environmental conditions as well as characteristics of the patient including cardiovascular function, skeletal muscle training and psychological factors (2).

1. Types of Exercise Encountered in Occupational Tasks

Occupational tasks require different types of exertion, performed singly and in combination, and vary in the number and mass of muscle groups involved. The major types of exercise usually performed are *dynamic* (isotonic) and *static* (isometric), or a combination of both dynamic and static exertion.