Smoking: a major predictor of left ventricular function after occlusion of the left anterior descending coronary artery

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SUMMARY  The major predictors of left ventricular function after coronary artery occlusion were assessed in 108 consecutive patients who had complete occlusion of the left anterior descending artery as the only important lesion demonstrated at angiography between June 1978 and June 1983. A scoring system was used to identify regional damage on left ventriculograms. Forty two patients were classified as having good left ventricular function and 66 as having varying degrees of impairment. Apart from a history of myocardial infarction, the only variables discriminating between those with good and those with impaired left ventricular function were the area of distribution of the artery beyond the occlusion and cigarette smoking. Hypertension, hypercholesterolaemia, family history of vascular disease, diabetes, obesity, duration of angina, age, and presence of identifiable collaterals were not discriminators. Smoking was itself significantly associated with a history of infarction; but after controlling for this, smoking exerted a significant additional effect on the amount of left ventricular damage.

It is concluded that smoking is not only a risk factor for myocardial infarction in patients with single left anterior descending artery occlusion, but that it is also a major factor in determining the extent of associated left ventricular damage.

Patients and methods

One hundred and eight consecutive patients with isolated left anterior descending occlusion were identified from the 2150 patients who underwent coronary arteriography in our laboratory between June 1978 and June 1983. All patients had complete occlusion of the left anterior descending coronary artery, with no important (> 50% diameter) stenoses in the circumflex or right coronary arteries. In all patients the left ventriculogram was technically adequate for assessment of regional wall motion.

The severity of left ventricular dysfunction and of coronary artery disease was assessed in each patient by the scoring system devised by Brandt et al. Representative systolic and diastolic frames from the 45° right anterior oblique left ventriculogram were traced and the ventricular outline in diastole was divided into five segments: the anterobasal, anterolateral, apical, diaphragmatic, and posterobasal. Depending on its systolic wall motion, each segment
was given a numerical value (normal = 0, hypokinetic = 1, akinetic = 2, dyskinetic = 3), and the scores of the five individual segments were summed to arrive at an overall left ventricular score. Patients with scores of 0–2 were arbitrarily classified as having "good" left ventricular function (group 1) and those with a score of 3 or more as having impaired left ventricular function (group 2).

Coronary arteriograms were performed in multiple views by the Sones' or Judkins' techniques. We used Brandt's scoring system\(^3\) to give each stenosis of a major coronary artery a numerical value that depended upon its site (proximal or distal), its severity, and its area of supply. In this system, possible coronary scores range from 0 (if all arteries are normal) to 15 (if there is a total occlusion of all three coronary arteries).

Identifiable collateral vessels were graded according to the degree of opacification of the distal left anterior descending artery seen during coronary angiography: zero if a distal vessel was not identified; 1 if the distal vessel was seen only faintly and incompletely; 2 if the distal vessel was completely opacified; and 3 if the distal segment was as well opacified as the vessel being injected. This assessment was made after both right and left coronary injections, the two values being summed to give a collateral score.

To determine the important predictors of left ventricular function after left anterior descending artery occlusion, we compared several pre-existing variables in the patients with good (group 1) or impaired (group 2) left ventricular function. These included smoking, hypertension, family history of premature vascular disease, serum cholesterol, diabetes, body mass index, and duration of angina, the latter indicating possible predisposition to collateral formation. We also assessed variables that presumably result from left anterior descending coronary artery occlusion—area of supply as reflected by coronary artery score, a history of infarction, and the presence of abnormal Q waves on the electrocardiogram.

Patients were classified according to smoking history at the time of infarction or first presentation as: non-smokers, patients who had not smoked for at least two years; and light smokers, those smoking <10 cigarettes a day and pipe and cigar smokers; moderate smokers, between 10 and 20 cigarettes daily; heavy smokers >20 cigarettes a day. A period of abstention of two years was chosen for non-smokers because data from the Framingham study, confirmed by Rosenberg \textit{et al}, showed that by this time the risk of a cardiovascular event reverted to that found in people who had never smoked.\(^4\)\(^5\)

Patients with a blood pressure >150 mm Hg systolic or >95 mm Hg diastolic or who had been treated for hypertension were regarded as having hypertension. Patients were classified as having a family history of ischaemic heart disease if ischaemic heart disease had developed in a first degree relative aged <60. Fasting serum cholesterol, height, weight, and body mass index were measured at the time of cardiac catheterisation.

We performed stepwise discriminant analysis (Statistical Package for the Social Sciences) of the variables shown in Table 1 to discriminate between group 1 (good left ventricular function) and group 2 (impaired left ventricular function) patients. This program shows variables that discriminate between the two groups and at each step in the analysis selects the variable that gives the best discrimination. A statistic which is used to evaluate a discriminating variable is termed "F to enter" and in this calculation any value >3.96 is significant at the 5% level. The same discriminant analysis was used when a history of infarction was included in the analysis.

**Results**

Of the 108 patients with isolated complete occlusion of the left anterior descending coronary artery, 90 were male (mean age 51 years, range 34–69 years) and 18 were female (mean age 51 years, range 34–63 years). Most had symptoms of angina (74), left ventricular failure (10), or arrhythmia (4) and were studied with a view to bypass surgery. Twenty were symptom free or had only slight symptoms; most of these were younger patients with a history of myocardial infarction or of atypical chest pain in whom angiography had been performed mainly as a guide to prognosis.

Ventricular function was normal (score of zero) in 22 patients and abnormal in 86. Figure 1 shows the distribution of ventricular scores among the 108 patients studied. Forty-two patients had ventricular scores of 0–2 and were included in group 1 ("good" left ventricular function). The remaining 66 patients had ventricular scores of 3–9 and were included in group 2 ("impaired" left ventricular function). The cut off point of 0–2 for group 1 was chosen arbitrarily because the statistical significance of the results was not different if group 1 patients included only those with scores of 0–1 or was extended to include those with scores of 0–3.

Figure 2 shows the coronary artery scores in the 108 patients ranged from 3 to 9.

Stepwise discriminant analysis of the pre-existing variables in Table 1 shows that only two, smoking (p < 0.0005) and left anterior descending coronary score (p = 0.0005) are significant discriminators. The program selects smoking classification as the
best discriminator. In group 1, 22 (52%) of 42 were non-smokers compared with 11 (17%) of 66 in group 2. After the coronary score is included the "F to enter" value of smoking remains relatively unaffected. Had there been a good association between the effects of these two variables on total left ventricular score, the fitting of one would have much reduced the "F to enter" of the other. There is, therefore, a largely separate effect of the two variables on left ventricular function as determined by total left ventricular score.

Table 2 shows the results when factors that presumably result from total left anterior coronary artery occlusion—history and electrocardiographic evidence of infarction and formation of angiographically visible collateral vessels—were included in the analysis. History of infarction is a significant discriminator (p < 0.00005) but not the presence of abnormal Q waves in the electrocardiogram (p = 0.28) or the formation of identifiable collateral vessels (p = 0.91). When history of infarction was included in the analysis, however, the "F to enter" value of smoking remained highly significant (p = 0.008); that is smoking remained an additional independent discriminator. Also discriminant analysis showed that smoking was significantly more common in those with a history of infarction (p = 0.014).
Smoking and left ventricular damage

Discussion

In this group of patients in whom complete occlusion of the left anterior descending coronary artery was the only important lesion ventricular function ranged from normal to severely impaired. The differences presumably reflect variations in the incidence and extent of myocardial infarction. Of the variables analysed in an attempt to explain these differences, apart from a history of myocardial infarction, only the area of supply distal to the left anterior descending occlusion and history of cigarette smoking were of significance in discriminating between patients with good and impaired left ventricular function. Discriminant analysis established that these two variables were acting independently. This remained true if the cut off point for group 1 ("good" ventricular function) was taken at a ventricular score of 2 or 3, and even if the nine light smokers in this study were classified as smokers rather than non-smokers. Age, sex, family history of vascular disease, or abnormal Q waves in the electrocardiogram, collateral score, or the presence of other major risk factors for coronary artery disease did not discriminate between the two patient groups.

It is known that lesions of the left anterior descending artery at different sites are associated with differences in left ventricular function. Patients with stenoses of the proximal left anterior descending artery have greater depression of ejection fraction, more abnormal wall motion, and more profound exercise dysfunction than do similar patients with distal stenoses. Such findings are not unexpected because the quantity of myocardium dependent upon a major coronary artery decreases as flow progresses more distally in the vessel. The results of the present study accord with these findings.

Cigarette smoking is associated with more severe and extensive coronary atherosclerosis and smokers have a considerably increased risk of both myocardial infarction (as shown here) and cardiovascular death. This increases with the number of cigarettes smoked. In this study we also found that after occlusion of a single major coronary artery left ventricular function was more depressed in heavy smokers than in a group of light and non-smokers with similar lesions. This association is independent of the site of arterial occlusion, the size of distribution of the artery, and a history of infarction and suggests that cigarette smoking independently affects the amount of myocardial infarction that results from the occlusion of a particular artery.

Support for this suggestion comes from a study of 3997 patients undergoing coronary angiography which showed a greater frequency of previous myocardial infarction in smokers than in non-smokers with coronary artery disease of comparable severity. Left ventricular function was not assessed in that study, however, and the inclusion of a heterogeneous group of patients with multivessel disease makes separation of the effects of smoking difficult. Epidemiological studies have also shown that although smoking is strongly related to myocardial infarction and death, the relation to angina pectoris is modest, if it exists at all. These findings also suggest an independent effect of cigarette smoking on the likelihood and size of myocardial infarction.

Acute thrombosis of a previously chronically narrowed coronary vessel is the pathogenic mechanism responsible for most transmural myocardial infarctions and cigarette smoking may promote thrombosis by increasing platelet aggregation and adherence. Cigarette smoking seems to affect the biochemical composition of atherosclerotic plaques and this may influence their rates of growth and thrombogenic potential. A recent study has also shown that production of prostacyclin, a potent vasodilator and platelet antiaggregatory agent produced by vascular endothelial cells, is depressed in smokers.

The importance of these smoking induced changes lies in their potential for altering the time course over which occlusion of a vessel takes place and consequently the amount of left ventricular damage that results. We suggest that cigarette smoking mainly affects ventricular function by promoting thrombus formation at sites of atherosclerotic narrowing, and that in smokers the development of sudden complete occlusion may be more likely than progressive gradual narrowing of non-critically narrowed vessels. The consequences for left ventricular function will be great because the time taken for a stenosis to develop has a major bearing on the development and size of collateral vessels in both animal and human studies and collateral flow is a major determinant of the amount of myocardial damage occurring after coronary occlusion. Our finding that there was no significant difference in collateral score between patients with good and impaired left ventricular function may simply reflect the fact that coronary angiography only visualises epicardial coronary arteries with diameters >100 μm. Most human collaterals are smaller than this and are subendocardial. Therefore, the number and size of collaterals identified at angiography may have only limited relevance to collateral flow.

Changes in heart rate, blood pressure, plasma catecholamines, and carboxyhaemoglobin concentrations are also seen in smokers and these provide further potential mechanisms through which cigarette smoking may influence infarct size. Other
variables such as autonomic tone and drug treatment at the time of infarction may also be important. Whatever the mechanisms, however, the present study identifies a clear-cut association between smoking and ventricular function among a group of symptomatic and asymptomatic patients with occlusion of the left anterior descending coronary artery. Furthermore, the power of the association we found strongly suggests that the smoking habit is an important independent determinant of left ventricular function in patients with ischaemic heart disease.

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References
