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<th>Responses of plasma concentrations of A type natriuretic peptide and B type natriuretic peptide to alacepril, an angiotensin-converting enzyme inhibitor, in patients with congestive heart failure [4]</th>
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Insulin resistance, high prevalence of diabetes, and cardiovascular risk in immigrant Asians

Sr.—We read with interest the findings of Dhawan et al., who highlighted the importance of hyperinsulinemia, central obesity, and physical inactivity as risk factors in Asians with angiographically significant coronary artery disease. Both British and Indian Asians were found to share a predisposition to insulin resistance and its associated metabolic abnormalities—and hence a high cardiovascular risk. They conclude that this finding is likely to favour a genetic rather than an environmental basis for the recognized high mortality in this ethnic group. However, migrants are not a random sample of the original population, and their “selection” is likely to be determined by several health and socioeconomic factors, which are likely to influence their morbidity and mortality. If this environmental effect were a significant determinant of cardiovascular risk, one would not expect to see the high mortality from ischaemic heart disease (IHD) that has been recorded in Asians in South Africa, Trinidad, and Singapore, who emigrated over a century ago.

We recently reported a survey in which we studied the cardiovascular risk factor profile of all Asian men admitted with acute myocardial infarction during 8 weeks to our city centre district general hospital in Birmingham, England, and to the San Fernando General Hospital, in Trinidad 4; 74 patients were studied (55 patients (mean (SEM) age 58±1 (1-4) in Trinidad (Trinidad group) and 19 in Birmingham (62±1 (2-6)) (UK Group) (table). We also found that mean systolic and diastolic blood pressures were higher in those with hypertension in the Trinidad group 10/2 (16-9) to 1/4 (11-4) mmHg than in the UK group (120-8 (25-4) to 75-0 (13-4) mmHg, P < 0-05). Though Asians in Trinidad have in many ways adapted to the lifestyle of the host population, this does not appear to have reduced their cardiovascular risk profile, because those admitted with acute myocardial infarction had, in fact, a greater prevalence of central obesity, smoking, and higher blood pressures than a similar group in England.

Central obesity and physical inactivity were common to both communities in England and Trinidad and their relation to insulin resistance may be particularly important in Asians with IHD. Our data support the hypothesis of a genetic predisposition to central obesity and diabetes in Asians that seems to have been retained by

### Quality of life in heart failure treated with enoximone

Sr.—The Enoximone Investigators concluded that in severe heart failure enoximone reduces survival but has a beneficial effect on the quality of life of survivors. As they indicate, if true, this is an important finding with interesting and difficult implications for drug regulatory bodies and for clinical practice. However, a statistical artefact could explain the result without there necessarily being any benefit from enoximone.

At randomisation, the two groups were reasonably well matched for measures of severity and prognosis. As the trial progressed there was differential mortality between the groups. It is not unreasonable to assume that those who died had more severe disease and a worse quality of life than those who survived. The even if there were no change at all in quality of life, the survivors will have a better average quality of life than the study group as a whole. Since more died in the enoximone group, this group will have lost more patients with poor quality of life, and enoximone will appear to have been beneficial. Other evidence from the paper suggests this interpretation of the results, namely, the greater proportion of enoximone treated patients withdrawn from the trial and the greater proportion admitted to hospital. The improvement seen in the disease-specific measure after 2 weeks (when deaths were 8/18, from table 3) was probably genuine, but the lack of differences at three months and one year was unlikely to be because of the power alone (although 95% confidence intervals for the differences in means were not quoted). Indeed, as one would have expected the group with more deaths to have a better average quality of life for the reasons given, the possibility arises that quality of life actually diminished on enoximone treatment.

Apart from the baseline comparison, we are not able to interpret the Nottingham Health Profile (NHP) data at all. For example, the median physical mobility score in the enoximone group was 22 at one year, but there is nothing to compare this with. We are not told what the median baseline score was for the placebo treated patients nor longer form an adequate control group (as they are no longer matched with the enoximone group). It would have been interesting to see the results on the other NHP dimensions.

There are ways round this problem, but none is fully satisfactory. We can assign all patients who died to the bottom half of the distribution of quality of life scores for example, by giving them all the worst possible score). The median is then not biased for comparative purposes. However, if more than half the subject sample or median becomes uninterpretable (although the Mann-Whitney U test remains valid). Alternatively, end points can be dichotomised into those alive with improved quality of life on one hand and those alive with those who are alive with worse quality of life on the other. These can be analysed as proportions, but with consequent loss of statistical power. Finally, changes from baseline can be quoted for survivors only. These can be interpreted as uncontrolled observations (a so-called “before and after” study). So long as the questionnaire has reasonable test-retest reliability, a beneficial treatment effect is the most likely explanation for any improvement seen, although a placebo effect cannot be excluded.

### Responses of plasma concentrations of A type natriuretic peptide and B type natriuretic peptide to alacprin, an angiotensin-converting enzyme inhibitor, in patients with congestive heart failure

Sr.—Yoshimura and colleagues showed that the response of plasma brain natriuretic peptide (BNP) after administration of the angiotensin-converting enzyme inhibitor, alacprin, occurred later and lasted longer than the response of plasma atrial natriuretic peptide (ANP), and that the changes in pulmonary capillary wedge pressure did not correlate with plasma BNP. 1 This lack of correlation may be the result of this difference in responses.

It may be relevant to recall that BNP has a longer half-life than ANP and that in BNP is bi-exponential, with a slow phase of about 20-7 minutes, as estimated by Nakao's group. 2 My own estimate of the slow half-life was 37-1 minutes, 1 and there may also be a qualitative difference in the synthesis and release of BNP.
Transcatheter occlusion of cardiac defects

Sin—Gatzoulis, Redington, and Rigby et al reported their experience with transcatheter occlusion of the ductus arteriosus and of atrial and ventricular septal defects with the Rashkind ductal and atrial septal occluder. We have considerable experience with this device in nearly 400 patients with atrial septal defects, 120 patients with a patent ductus arteriosus (PDA), and eight patients with ventricular septal defects (unpublished observations).

Patent ductus arteriosus—Our experience with the buttoned device indicates that it has several advantages over the Rashkind device. One advantage is that it can be closed in a smaller size and in a patient with a small ductus by adjusting the clip. In all infants and children we were able to implant the device through a 7F delivery sheath rather than 8F and 11F sheaths, which have been used in most of Gatzoulis's patients. The success rate for device implantation was 86% for the Rashkind device and 97.5% for the buttoned device. Problems such as embolisation into the left pulmonary artery, inability to occlude the defect because of the dimensions of the heart, and severe haemolytic anaemia reported by Gatzoulis et al were not seen in our 120 patients. Residual shunts on colour flow mapping, particularly in infants who had 17 mm Rashkind umbilical devices implanted were higher than expected, but these shunts were not present when we saw the buttoned device. Finally, there has been increasing concern about the development of stenosis of the left pulmonary artery after implantation of the Rashkind device, especially in younger children. Thus the buttoned device seems to have several advantages over the Rashkind device and it is hoped that, with further clinical trials, the buttoned device will prove to be useful in transcatheter occlusion of arterial ducts of all types and sizes.

Atrial septal defect—Redington and Rigby modified the Rashkind and used it to close interatrial communications of various types. Of the 11 patients with fenestrated Fontan, there were two (18%) procedural failures. They were successful in closing defects in 15 patients, with two residual shunts of less than 1mm Hg with left-to-right shunts; the two remaining patients required surgical removal of the device and closure of the defect. In our more recently reported experience, which analysed the data of the first 180 patients, 14 devices (7·7%) were dislodged. The dislodgement rate has improved with experience and with successive generations of the device. In the first and second generation devices became dislodged in about 11·5% of patients. However, in the third generation device the dislodgement rate was 3·1% (2 of 65); this has further decreased to less than 1% in the fourth generation buttoned device (unpublished observations). Redington and Rigby placed a bend in the arm of the device. Such a bend is thought to be the reason why the arms of the Rashkind device break. We believe the device was withdrawn from clinical trials.

Ventricular septal defect—Rigby and Redington concluded that their data do not support the routine use of a Rashkind PDA occluder to close subaortic or subpulmonary septal defects. We agree. Our own experience in occluding ventricular septal defect with buttoned device, though successful, is limited. Therefore, we cannot draw definitive conclusions on the superiority of the buttoned device.

In conclusion, the reports of Gatzoulis, Redington and Rigby show the usefulness of the Rashkind PDA occluder in highly selected patient subgroups. We submit that the buttoned device has a greater utility in a wider range of patient subsets, although definite conclusions can only be drawn after longer clinical trials.