

whom 8% developed major depressive illness during the year before the interview.¹² After controlling for gender and family history of depression multivariate analysis indicated the importance of both job factors (lack of intrinsic rewards, conflict at work, and unclear job responsibilities) and home factors (marital strain and psychiatric disorder in spouse). Of the work environment the authors wrote, "Prevention and lasting remediation are most likely to result from bringing about positive changes in the cultural patterns characterising the business world. Such changes might attempt to strike a better balance between narrowly defined economic interests of the firm and broader humanistic concern for the mental and emotional needs of workers and their families."

With the current changes occurring in the NHS it is appropriate to consider whether similar factors affect its workforce—2.1 million days are apparently lost each year through mental disorders.¹ Radiographers, dentists, nurses, and doctors all feature in the top 14 jobs with a raised standardised mortality for suicide. It is less well recognised that the standardised mortality ratio for doctors' spouses is 275.⁵ Most other research has concerned the doctors themselves.

Surveys with the general health questionnaire have shown alarmingly high scores among junior house officers: 50% showed emotional distress, 28% had scores indicating possible depressive illness, and 6% had thoughts of suicide.¹³ The stressors most closely correlated with these symptoms included overwork, effect of work on personal life, relationship with consultant, and clinical decision making.

Stresses documented in general practitioners include night calls, emergencies during surgery hours, and interruption of family life.¹⁴ In this context clearly the separation of "job" and "home" stressors is artificial, and future studies need a comprehensive assessment of both.

Fortunately, the US employee assistance programmes and a recent scheme in the Post Office have shown the financial benefits of providing help for those experiencing stress at work, which should encourage the adoption of such schemes by large employers such as the NHS.¹ The recent research commissioned by the Department of Health on stress and the NHS workforce should identify more clearly than hitherto the job stresses at all employment grades, with a view to identifying where the work environment needs to be changed and where the adoption of employee assistance schemes might be most cost effective.

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"Self referral": a potential conflict of interest

The GMC should produce tighter guidelines

The break up of the NHS into discrete independent units has produced more entrepreneurial activity in an attempt to increase patient flows and "profit." A government that encourages commercialism in medicine should not be surprised to see comments in the medical press such as "Shares in laser centre selling fast"¹ and comments like "GP fundholders are channelling 43% of their inpatient work through their own limited companies."²

In private practice the relation between an episode of care and the fee that it attracts should be crystal clear to the patient. But this may become obscured when a doctor owns part of a facility and refers patients to it, thus increasing profit, without the patient being aware of a potential conflict of interest. Experience of this "self referral" is greater in the United States. In Florida one study showed that over 40% of doctors directly providing care had financial interests in health care businesses to which they could refer their patients for services, a large proportion of these being diagnostic imaging centres.³

Self referral significantly increases the costs of health care. A Californian study showed that doctors who referred patients to facilities that they owned initiated physical treatments 2.3 times more often than doctors who referred patients to independent facilities; for the self referral group

the total cost of psychiatric evaluation per case was 26% higher. Of magnetic resonance imaging scans requested by self referring doctors, 38% were clinically inappropriate compared with 28% of scans requested by doctors without a financial stake in the imaging centre.⁴ Another American study found that doctors who used imaging equipment in their offices obtained investigations 4.5 times more often than doctors referring through a radiologist and, as self referring doctors charged more, the total bill was up to 7.5 times higher.⁵

Responding to this and other information,⁶⁻⁹ the council on ethical and judicial affairs of the American Medical Association has published detailed guidelines.¹⁰ These state that doctors should disclose any relevant financial interests to patients when referring them. Patients should be given a list of alternative facilities and be assured that choosing one in which the doctor has no financial interest will not affect their medical care. Financial interests also have to be disclosed to a third party payer.

In Britain the General Medical Council has stated that "where doctors have a financial interest in an organisation to which they propose to refer a patient for admission or treatment, whether by reason of a capital investment or a remunerative position, they should always disclose that they

have such an interest before making the referral."¹¹ Does this happen in practice?

Occasions arise when doctors can provide privately services that the NHS cannot fund, but comprehensive guidelines are needed for these because of the ethical issues that they raise. In view of the changing nature of medical practice in Britain the General Medical Council should assess the effectiveness of its current recommendation and consider updating it along the lines of the American model.

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Does cimetidine cause weight loss?

Confounded expectations result in a conflict of evidence that is simply baffling

The ideal topic for a young researcher is one in which recent publications in peer reviewed journals have come to mutually incompatible conclusions. The tyro researcher is then presented with a ready made protocol and an expectation that a replication of the study will support paper A and refute B, or vice versa, or (more probably) reach an intermediate conclusion which throws some light on the reason for the conflict in results. Such a situation arises in this issue of the *BMJ*.

Paper A, by Støa-Birketvedt on p 1091, reported the outcome in 30 overweight subjects given 200 mg cimetidine as a suspension three times a day 30 minutes before meals, together with a fibre supplemented diet designed to supply 5MJ (1200 kcal)/day and 30 well matched controls given the same diet but a placebo suspension. The mean (SD) weight loss over the next eight weeks was 9.5 (2.1) kg in the treated group but only 2.2 (1.3) kg among the controls.¹ Paper B, however, by Rasmussen *et al* on p 1093, reports a replication of the study in paper A, in which the authors found no significant difference in weight loss between the cimetidine and placebo groups (5.7 kg and 5.9 kg respectively).² Paper B suggests that the results in A may have arisen because the subjects were not really blind to the medication taken: this may be so, but it does not resolve the problem. A mean loss of 9.5 kg in eight weeks is a remarkably good result, and if such weight loss could be achieved by simply telling patients they were on cimetidine that would be a therapeutic triumph.

In this situation an experienced editor may suspect that there is something wrong with the experimental data, but Dr Støa-Birketvedt has been commendably frank and generous in making the raw data available for examination. (The data and the patients have also been re-examined by a committee at Oslo University and confirmed to be accurate.) The results are correctly calculated and do not arise as a result of selective attrition. Papers about the treatment of obesity often fail to take account of the fact that the mean weight given for a group of patients at different points in time may refer to an ever diminishing cohort. As less successful weight losers drop out

there is a false appearance of continuing weight loss in the group. Paper A started and finished with 30 patients in each group, so selective attrition cannot explain these findings.

The individual data on which paper A is based show a remarkable uniformity of weight loss between individuals in each treatment group and also within individuals from week to week. Both features are hard to explain. Among groups of overweight people weight loss is usually more rapid in the first week of dieting than subsequently, probably because naive dieters lose glycogen stores with associated water first during the dieting period, and more energy dense fatty tissue later.³ This effect is seen in the results of paper B, but not paper A, in which the mean weight loss was the same in the first two weeks of cimetidine as in the last two weeks. Another expected effect not seen in the results of paper A is that individuals with a higher initial weight show a greater weight loss than those with a lower initial weight. The mean weight loss in the five heaviest people, who initially weighed more than 90 kg, was only 0.5 kg greater than that in people who initially weighed less than 70 kg, whereas a difference of at least 2 kg in weight loss would be expected. Finally, the absolute weight loss among the cimetidine group in paper A is surprisingly high, and that of the control group surprisingly low, while the mean 5.8 kg loss in eight weeks in paper B is what might be expected in well supervised subjects with an initial body mass index (weight (kg)/height (m)²) of 34 and a diet designed to supply 5MJ/day. It remains baffling that cimetidine and placebo should have such different effects in Norway and Denmark.

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