council of the Royal College of General Practitioners and the General Medical Services Committee (Wales).

In overall terms our deputation feared that the general principles of the referral system were compromised, access to specialist psychiatric services was ill defined, and the declared role of the mental health team in primary care could lead only to fragmentation and confusion. Furthermore, the contractual obligations of the general practitioner were totally bypassed.

The deputation received a sympathetic hearing, and it was with great disappointment that we read the final paper, *Mental Health Services, a Strategy for Wales*, issued in June 1989. Little has changed from that set out in the consultation document, and we are convinced that if the recommendations of this paper are implemented the task of treating psychiatric disorder in Wales is likely to be muddled and expensive.

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Rape and subsequent seroconversion to HIV

SIR,—The paper by Dr S Murphy and colleagues highlights the potential risks and worries for women who have been sexually assaulted during the current phase of the HIV epidemic.¹ If we assume that such assailants are heterosexual men the overall risk of transmission is likely to be small in view of the present low carriage rate for the virus among this group in Britain.¹ In addition, vaginal intercourse is possibly a less likely mode of transmission than anal intercourse.³ The situation for male victims of sexual assault may, however, be different.

Male "rape" as a legal term does not exist in the United Kingdom, as rape specifies forcible vaginal penetration. Furthermore, male sexual assault is not recognised as a distinct entity within the 1976 Sexual Offences (Amendment) Act.⁴ This may well have consequences for the number of cases reported and for public awareness of such events.⁵ Documented cases of sexual assault of men by women have been reported,⁶ but it is generally thought that men are more commonly assaulted by other men.

In the United States it has been estimated that only 10-20% of all sexual assaults are ever reported.⁷ In addition, there is some evidence to suggest that male victims are more reluctant to come forward than their female counterparts, possibly related to the fear of being labelled as homosexual⁸ or to society's concept that a "real man" cannot be raped.⁷

Partly as a consequence of its legal non-existence there are no reliable figures of the frequency and nature of male sexual assault in Britain. The only organisation providing care for victims of male sexual assault in Britain at present is "Survivors," a service run by volunteers to which various government and charitable agencies refer their clients. The service provides a telephone help line (currently answering just over 100 inquiries a month) together with counselling and support. No studies to date have assessed the risk of sexually acquired infection in male victims of assault,⁹ but of 512 victims known to Survivors in 1988, 24 of 73 who sought medical advice were diagnosed as having a sexually transmitted disease presumed be consequent on their assault. Data on HIV seroconversion are not available, but 148 of the 512 victims reported skin or mucosal bleeding, and 278

expressed concern about the possibility of transmission of HIV. In 92 of the cases the assailant deliberately threatened the victim with the possibility of contracting HIV infection as a consequence of the assault.

It has been suggested that between 50%¹⁰ and 82%⁴ of assailants of male victims are either homosexual or bisexual. The assailants are therefore in relatively higher risk groups for HIV infection than heterosexual assailants of women. Furthermore, anal penetration, bloody non-genital violence, and multiple assailants are more likely when the victim is male.⁷

Rape treatment centres have been set up primarily for female victims and may lack the skills to deal with men. We believe that an increased level of awareness of male sexual assault is needed among the general public and especially by health care professionals to encourage victims to come forward. Only when this happens can the scale of the problem be fully grasped and appropriate treatment provided.

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Passive smoking and cardiorespiratory health in Scotland

SIR,—Mr Peter N Lee¹ implies that our observation of increased risk for four respiratory symptoms and two cardiovascular symptoms, mortality from lung cancer, mortality from ischaemic heart disease, all causes of death related to smoking, and mortality from all causes in passive smokers compared with controls² can be explained by bias—that of smokers declaring themselves to be lifelong non-smokers. He cites (presumably) lung cancer, for which if, as he supposes, no increased risk is associated with exposure to environmental tobacco smoke and the "true" risk is increased 20 times in active smokers,

2% of smokers denying smoking would result in observed relative risks of 1.74 for passive smokers and 6.90 for active smokers. This is illustrated in his table, using the distribution of subjects in the smoking groups defined in our study. But why does he use only men and ignore women when our analysis and results were based on both sexes and women comprised 84.2% of our passive smokers? If he had included women 5.0% of passive smokers and 3.9% of controls would have smoked (table I)—quite different from the figures of 15.6% and 6.8% respectively presented in his table. Our figures in turn produce an observed relative risk of 1.12 for passive smoking (substantially less than the 1.74 he quotes by selecting only men) and considerably less than our study finding of 2.41 for lung cancer among passive smokers. Clearly, misclassification owing to the 2% rate of denial of smoking he suggests does not explain our finding.

In addition, higher rates of denial of smoking do not produce sufficient bias to explain our risk for lung cancer. Table II presents the effect on the basis of a "true" relative risk of 20 for active smoking and 1 for passive smoking and of rates of denial varying from 1% to 10%. Two facts emerge: firstly, the bias in the relative risk for passive smokers does not increase linearly as the rate of denial increases, it flattens considerably; secondly, the observed relative risk for active smokers diminishes rapidly as the rate of denial increases.

TABLE II—Relative risks for passive and active smoking for varying rates of smoking

Rate of denial (%)	Observed relative risk	
	Passive smokers	Active smokers
1	1.07	13.95
2	1.12	10.67
3	1.15	8.61
4	1.18	7.17
5	1.19	6.14
6	1.20	5.35
7	1.21	4.73
8	1.22	4.23
9	1.23	3.82
10	1.23	3.47

This is important, as by comparing the observed relative risk for active smokers from the table with the relative risk found in the study an upper bound can be defined for the rate of denial. The study relative risk for active smokers compared with lifelong non-smokers is 5.85. This would be incompatible with rates of denial greater than 5%. Therefore, the largest relative risk to be expected among passive smokers due to this form of bias when the "true" risk is unity is 1.20. A relative risk of 2.41 was found in our study.

Again, the same approach applied to ischaemic heart disease assuming a "true" relative risk of 3 for active smokers and 1 for passive smokers and a rate of denial of smoking of 5% produces an observed relative risk of 1.05 for passive smokers and 2.42 for active smokers. Thus if the relative risk for active smokers is considerably less than 20, as in all the conditions we considered other than lung cancer, the effect of misclassification is to produce only small biases in the relative risk for passive smokers. Our risks for each of the respiratory symptoms, cardiovascular symptoms, and cate-

TABLE I—Differential misclassification caused by 2% of index subjects denying smoking regardless of cohabitee's smoking habits

Exposure group*	Observed distribution†	"True" distribution	Effects of denial	Percentage who have smoked
Controls	917	881	+36	3.9
Passive smokers	1538	1461	+77	5.0
Single smokers	1751	1787	-36	
Double smokers	3791	3868	-77	

*As defined by Hole *et al.*†Data from table I of Hole *et al.*

gories of mortality quoted are well in excess of that produced by the form of bias suggested. Also, as control subjects experience some level of environmental tobacco smoke² our estimates of risk could be conservative.

Mr Lee misunderstands our use of urinary cotinine concentrations in passive smokers. We were using published data to establish whether our study had sufficient statistical power to detect the size of risk that might be expected among passive smokers. If Mr Lee is correct and urinary cotinine concentrations are equivalent to a lower dose than assumed then our decision not to rely solely on statistical significance as evidence of a genuine effect was definitely correct. Ours was a cohort study of a general population and was not subject to the biases associated with a case-control design. In addition, subjects reported their own smoking histories, and environmental exposure was based on record linkage of cohabitants, thereby avoiding the need to rely on self reporting of passive exposure.

Our observations on lung cancer may be based on only nine deaths but are consistent with the result of a meta-analysis³ combining 13 separate studies, which concluded that breathing other people's tobacco smoke causes lung cancer. The importance of our study lies in the estimates of risk for ischaemic heart disease (based on 84 deaths), all causes of death related to smoking (175 deaths), mortality from all causes (263 deaths), respiratory symptoms (292 cases), and cardiovascular symptoms (117 cases). The consistent increase in risks for such a wide variety of health outcomes from an unbiased prospective cohort study together with a dose-response relation in passive smokers strongly suggests that there is now a case to be answered against passive smoking that extends beyond the causation of lung cancer.

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Referrals from general practice to hospital outpatient departments

SIR,—One aspect highlighted in the report by Drs John Emmanuel and Nigel Walker¹ is treatment of skin disorders in general practice. Proposals in the white paper are likely to encourage more minor surgery to be undertaken by general practitioners. This may be more cost effective (although our own experience indicates that this may not necessarily be so), but skin surgery should be undertaken in general practice only if the diagnosis is certain—otherwise referrals may be increased rather than decreased as intended. We report two problems that resulted from inappropriate skin surgery in general practice.

A 49 year old woman had a pigmented lesion removed by curettage and cautery from her lower leg by her general practitioner. Histology showed malignant melanoma, but it was impossible to ascertain the depth of the tumour on the basis of the inadequately thin curettage specimen. The patient then had a wide excision and graft, but it is

possible that she would not have required an extensive operation because narrow excision margins can sometimes be adequate for very thin melanomas.

In another patient, a 46 year old woman, a slightly raised nodule on the leg was treated by curettage and cautery by her general practitioner. Histology showed invasive squamous cell carcinoma and the patient was referred for further advice. Because it was difficult to know the adequacy of the initial treatment the patient was committed to prolonged follow up to exclude recurrence of the lesion.

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Provision of services

SIR,—It seems to be the custom that when a specialist advisory committee pronounces on how services should be provided this is accepted, but there are occasions when someone needs to stand up and say "You are wrong."

The North West Thames ear, nose, and throat regional advisory subcommittee says that inpatient ear, nose, and throat services should be provided only in subregional specialist centres and not in the smaller district general hospitals. I have been the anaesthetist for three to four ear, nose, and throat lists per week for over 20 years and know that most of these operations are everyday bread and butter surgery and that over half are on children. Indeed the commonest paediatric operations are ear, nose, and throat—tonsils, glue ears, etc. These services have always been available at the local hospital and to say they should all go to subregional centres is tantamount to saying all hernias and ingrowing toenails should go to specialised units. Not only does this deprive patients of what I would call a core service but it has profound knock on effects on most other services in the district general hospital through the possible loss of recognition of anaesthetic jobs. Before someone brings out the old chestnut of "Make rotations" I will answer "Just you try to."

We are facing this situation in North West Hertfordshire District, where the loss of inpatient ear, nose, and throat services will disadvantage our patients and could cause havoc with the hospital services as a whole. I am afraid that this may be only the beginning of specialist groups building their own little empires without regard to the patients and hospitals from whom they withdraw their services.

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Psychiatric illness among the homeless

SIR,—Dr Max Marshall describes a high proportion of residents of Oxford hostels for the homeless as being "long term psychiatric patients" and implies that they are deinstitutionalised long stay patients.¹ Our findings, however, suggest that hostel residents with psychiatric disabilities may have had numerous yet relatively brief hospital admissions and include those sometimes referred to as "revolving door" patients.

We are currently evaluating a psychiatric liaison

service to residents of a direct access hostel for homeless women in central London. Of 33 women seen to date, 26 are known to have had at least one previous psychiatric admission, but only four have spent periods of more than one year continuously as inpatients. We believe the current emphasis on deinstitutionalised long stay patients is misplaced: it is the needs of those with chronic, severe psychiatric disabilities in the community and the revolving door patients that are not being addressed. Deferring the closure of psychiatric hospitals² will have little impact on this large group of people. The Department of Health has stated that the forthcoming white paper on community care will contain plans to prevent the unplanned discharge of long stay patients into the community. These safeguards will be of no value to most severely disabled psychiatric patients in the community.

Dr Marshall's findings and our own data both show high levels of unmet need and are in keeping with most surveys of people with psychiatric disorders in the community. These findings clearly indicate inadequate provision of care, but they should not be used as evidence of the ineffectiveness of deinstitutionalisation programmes or properly planned and funded community services. The few controlled studies of selected patients discharged within carefully planned community programmes³ show that long term psychiatric patients (whether or not they have had long stay psychiatric admissions) can be maintained outside hospital without the deterioration in symptoms, poor psychosocial functioning, and readmissions that are all too commonly found in the surveys. Perhaps more importantly, the controlled studies in which patients' wishes and satisfaction have been recorded clearly show that they prefer to be treated in the community.

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Safety of Picolax in inflammatory bowel disease

SIR,—In view of the suggestion of Dr A J G McDonagh and colleagues that further evaluation of Picolax is merited¹ we would like to report our own experience with this preparation in a large cohort of children undergoing fiberoptic colonoscopy at St Bartholomew's Hospital. Between 1982 and 1988 we performed 534 colonoscopies on 412 children attending this hospital and, with few exceptions, Picolax was used routinely to prepare the colon before endoscopy. This included the 287 procedures performed on children with chronic inflammatory bowel disease (163 with Crohn's disease, 101 with ulcerative colitis, 23 with indeterminate colitis) that was either known to pre-exist or suspected and confirmed at the time of endoscopy. We found the preparation to be successful for cleansing the bowel and free of major complications.

Based on our experience we have developed the following regimen for preparing the colon before endoscopy in children. The child is given only fluids for 24 hours before the procedure and is given two doses of Picolax, one about 15 hours before endoscopy and the other three hours before. The dose is age dependent: children over 6 years