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appealing hypothesis that early postoperative luminal nutrition might have a beneficial effect on the function of the intestinal barrier in respect of permeability, bacterial translocation, and the subsequent development of septic complications has no supporting evidence at present.

What impact could the findings of this systematic review have on daily surgical practice? The review shows that there is no clinical benefit to starving patients in the early postoperative period after gastrointestinal resection. Further, the finding that postoperative infections can be reduced and hospital stay shortened by starting early postoperative enteral nutrition should challenge clinicians to consider this treatment. The findings pave the way for an appropriate multicentred trial to assess early enteral feeding in patients undergoing elective gastrointestinal resection. The patients recruited to such a trial should be stratified by nutritional status and type of surgical procedure. The outcome measures should include not just effects on wound infection, other infectious complications, and dehiscence of the anastomosis but also surgical fatigue, muscle function, quality of life after discharge from hospital, and cost effectiveness.

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**Equity versus efficiency: a dilemma for the NHS**

*If the NHS is serious about equity it must offer guidance when principles conflict*

Concerns about equitable provision and financing of health care have characterised the NHS since its foundation. Evidence of persisting and, in some cases, widening health inequalities, gathered since the publication of the Black report, has progressively raised equity to a high rank among health policy objectives. Though the general aim of reducing health inequalities appears uncontroversial, the practical notions of equity that should inform policy and the ways in which these should be implemented are far from clear. Even more importantly, there is no consensus on how to deal with policies that may cause a conflict between the goals of equity and efficiency—that is, those that may improve efficiency while increasing health inequalities or equity and efficiency—that is, those that may improve fairness while decreasing efficiency. The equity versus efficiency dilemma has been virtually ignored in the political debate, often leading to inconsistent judgments in the development of health policies.

In a report recently published by the NHS Health Technology Assessment programme we examined examples of the equity-efficiency dilemma that the NHS is facing. The analysis of three case studies—cervical cancer screening, renal transplantation, and neonatal screening for sickle cell disease—shows inconsistencies between NHS policies and a lack of guiding principles to support the pursuit of equity in health care.

The NHS policy on cervical cancer screening has been primarily aimed at maximising coverage by using powerful economic incentives to general practitioners. The issue of low participation by women at high risk (particularly those in disadvantaged socioeconomic groups) has been less of a concern. The programme could have achieved the same cost effectiveness with less extensive but more even coverage. The number of cases of invasive cancer avoided in 1997 is likely to be 60-85% of the number of cases that might have been avoided if screening rates had increased uniformly in different social groups after the introduction of target payments to general practitioners."
efficiency grounds, but—of more relevance for our purposes—not even on equity grounds, as some studies have shown that the public would rank older children over younger ones. Although explicitly formulated in some respects, this NHS policy again appears to lack a clear reference to a guiding equity principle.

Sickle cell disease disproportionately affects certain ethnic minority groups. The UK Standing Medical Advisory Committee recommended the use of universal, rather than selective, neonatal screening policies when ethnic minorities with a high risk comprise more than 15% of the population. At this threshold the cost of universal screening is as high as £430,000 to £1m per life year saved (depending on the ethnic minority mix) compared with selective screening. The adoption of universal screening does not appear to be justified by concerns for equity across ethnic groups, as the benefits to the white northern European majority would still be very small. Rather, it aims at reducing the number of cases missed because of inaccuracies in the selection. This NHS policy may reflect an aspiration to equal access for equal need, but one pursued at a very high cost. Significant efficiency gains may be sacrificed for what seems to be an inappropriate conception of equity in this context.

More examples of inconsistency can be found among current NHS policies, and even greater variation could be unveiled. But is it realistic to expect health policymakers to develop sound and consistent policies in the absence of evidence about the distributional effects of healthcare provision? Is it realistic to expect them to address the equity versus efficiency dilemma? A systematic review of the literature on healthcare economic evaluations published in 1987–97 shows a complete neglect of the equity dimension within the studies surveyed. Not only did these studies fail to incorporate equity measures in their cost effectiveness calculations, they did not even provide enough information for decision makers to make their own judgments about the distributional impact of given policies—for example, on the characteristics of the population affected by the policy or on the policy’s effectiveness and cost effectiveness in subgroups.

Our three case studies show the lack of a clear and consistent definition of equity and the failure to strike an acceptable balance between the policy goals of equity and efficiency when these conflict. In different ways researchers and policymakers share responsibility for the inconsistent pursuit of equity in the NHS.

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Screening for prostate cancer in the UK

Seems to be creeping in by the back door

Screening for prostate cancer is controversial. Findings from systematic and other reviews consistently conclude that there is insufficient evidence to recommend its introduction because of concerns that it may not improve survival or quality of life and may thus cause more harm than good. Current government policy in the United Kingdom, expressed in the NHS prostate cancer programme, confirms this view, but adds that “any man considering a PSA [prostate specific antigen] test will be given detailed information to enable him to make an informed choice about whether to proceed with a test or not.” This implies that asymptomatic men may have the test if they want, so there is now ambiguity about whether screening is supported and confusion about what this policy means in practice.

The assumption may be that most men will not want to be tested once they are informed of the uncertainties. In the United States several studies have shown that informed choice can reduce prostate specific antigen testing in some groups by up to one half. But this may not apply in the United Kingdom. A systematic review of the use of decision aids has shown that though such aids result in higher levels of knowledge, they have variable effects on the decisions themselves, with reduced preferences for prostate specific antigen testing found in two studies but no effect in two others. Further, close inspection of the landmark study shows that though prostate specific antigen testing was reduced by half among scheduled clinic attendees who viewed a video, a parallel (rarely quoted) trial found that only 3 out of 206 men attending free prostate spe-