MMR vaccination and autism 1998

There is no causal link between MMR vaccine and autism

Editor—Nichol et al considerably underestimate their case when they write “chance alone dictates that some cases [of autism] will appear shortly after vaccination.” Such a temporal association is unremarkable, given the epidemiology of autism and MMR vaccine.

Over the time described by Wakefield et al1 MMR vaccine was given to around 600 000 children each year in Britain2 and the prevalence of autistic spectrum disorders was 91/100 000.3

Assuming that the diagnosis of autism is evenly distributed over the second and third years of life and that the incidence over this period approaches to the current prevalence,4 over the eight years that the reported cases represent autism would have been diagnosed in around 364 cases in the two months after MMR vaccination (the time that the authors regard as noteworthy—((91/100 000) × 600 000) × 8) × (2/24) = 364). The reported cases therefore represent a fraction of the cases of autism whose onset coincides with the administration of MMR vaccine.

R Roberts Consultant in public health medicine North West Health Authority, Preswyfyl, Mold, Flintshire CH7 1PZ

Those giving MMR vaccine had no input into editorial

Editor—The editorial by Nicoll et al about a possible link between measles, mumps, and rubella (MMR) vaccine and autism1 is not helpful for those working in primary care who actually give the vaccine, are concerned about its potential and actual side effects, and have to deal with the effects of a scare story, as in this case. General practitioners, health visitors, and practice nurses, who all give this vaccine regularly, have had no formal input into the editorial and have not participated in the expert committee set up to review the MMR reports.

It is all very well to ask an expert committee for an opinion based on research in secondary, and even tertiary, care. A more valuable opinion of what really happens in practice, however, would be available from those who actually have the day to day experience of this vaccine in use, and that includes parents.

Public concern about the safety of MMR vaccine has not only been raised by the study of Wakefield et al1 in 1992 and 1993 one of the commonly used MMR vaccines was withdrawn because of recurrent reports of adverse effects. I remember seeing three cases of encephalitis after MMR vaccinations in my own baby clinic at that time, although thankfully these resolved.

Concerns after a television advertisement for the MR booster campaign in 1994 and 1995 left parents wondering whether the advertisement was meant to scare them into having the vaccination done. This may explain in part the media response to Wakefield’s study.5

What we would have needed from the editorial were some straightforward figures to help parents understand why their child should have this vaccine. An unvaccinated child is 20 times more likely to have a serious complication from measles, mumps, or rubella than he or she would be after MMR vaccination.

Finally, I assume that the last paragraph of the editorial contains an error: “While vaccine can be guaranteed to be without any risk …” No vaccine or drug can ever be guaranteed to be without any risk.

M R Klin General practitioner Rosendale Surgery, London SE21 8EZ

Medic al practitioners need to give more than reassurance

Editor—Can I present a challenge to Nicoll et al2 As a parent of young children I have become aware of a website—“The informed parent” (http://www.ucnc.edu/~aphillip/ WNW/vaccine/dvm.txt)—that is having a profound influence on many of my non-medical friends who have small children. Although the website is written by a non-physician, it is written in the style of a medical journal, which lends it more authority than it may merit. This website argues that vaccination is dangerous and unnecessary.

The issue is about risk and the perception of risk. There may also be a perception of secrecy about problems with vaccines in the United Kingdom, which is why this website is now so influential. In a consumer led NHS it is no longer sufficient for general practitioners and practice nurses to give simple reassurance.

The challenge is to produce information for parents that is accessible (including on the internet) and balanced, and that addresses the specific concerns that websites such as “The informed parent” engender. This is a particular example of patient empowerment, and the medical profession needs to do more than just convince itself of the safety of vaccination.

J Selway Senior registrar in public health Edensor, Tunbridge Wells, Kent TN1 3QL

The health of Gypsies

Problem of caring for travellers is British, not just European

Editor—The editorial by McKee1 on Gypsies, or Romas, was an interesting factual account of their origins as well as of the disadvantages and oppression they face as a minority group in continental Europe. Yet it is remarkable in that there is no mention of the disadvantages and oppression endured by the same minority community in the United Kingdom—from the Romans of folklore in the south of England to the tinkers of Scotland and to the New Age wanderers who seem to seek the romantic image without the responsibilities. Now they travel in trucks towing large caravans; when parked they are usually seen in lay-bys or on waste ground with lots of dogs, children, and rubbish.

In these unofficial caravan sites they live in the squalor and type of conditions prevailing in this country a couple of centuries ago—no clean or adequate water supply, no sewage disposal system, no rubbish collection, and certainly totally inadequate education, immunisation, and medical attention. The reaction is often to “get rid of them,” preferably by moving them on to some rundown housing estate in the next town but one. Local authorities have a duty to provide proper sites for travelling people, although this is too often frustrated by the “not in my back yard” (“nimby”) attitude and an inability of local authorities to attempt to work with the Gypsy culture by cooperating with their lifestyle.
It is easy to be disapproving about the unwelcome refugees in and around Dover. As far as our own travellers are concerned, it is a sad reflection on our society, and perhaps on the BMJ, that we behave as if nothing needs to be done in this country although it seems to be agreed that much needs to be done across the channel.

G A C Binnie  General practitioner, retired
Larkirk, Norham, Berwick upon Tweed TD15 1XL

Governments and Roma communities must help to improve outlook for Gypsies

Erratum—In his editorial on the health of Gypsies—an important minority in central and eastern Europe—McKee states that “health policymakers and researchers have paid little attention to the health needs of Roma people.”2 This is not true. The communist regime in the former Czechoslovakia spent considerable sums to improve the economic, health, and housing situation of the “proletarian” Roma population. Infant mortality of Gypsies decreased considerably and the number of Gypsies living in Slovakia more than trebled from the end of the second world war to 1990.2
Health care as well as the educational system in Slovakia was free and equally available to all Slovaks, Hungarians, and Gypsies. After the change from a socialist economy to a free market economy, unemployment among the Roma minority rapidly increased. The main reason was not racism but the low educational level and low working activity of Gypsies.

In 1996 we performed sociological research in various parts of Slovakia by a Gallup method. The table summarises some data from a representative sample of 1016 men aged 25–55 living in the multiethnic district of Levice on the border with Hungary (about 65% Slovaks, 30% Hungarians, and 5% Gypsies). The data are similar for both Slovaks and Hungarians and very different for Roma men: the Roma minority had an extremely high consumption of cigarettes, beer, and spirits; low consumption of milk, fruit, and vegetables; a lower cultural level; a higher prevalence of sleep disorders; a lower health status; and a higher birth rate.

Although Roma people continue to exist on the margins of society and their life expectancy is low, the number of Gypsies in Slovakia increased by 1.3% a year between 1991 and 1996.3 The reason for the high reproductive activity in Roma people is an economic one—regular financial support from state sources for each child. If this trend does not change, the Roma minority in 2010 will form a fifth of the young and middle aged population in Slovakia. Most Gypsies will have only elementary education and will be unemployed, chronically ill, and dependent on financial support from the state. A huge increase in crime could be expected, and the final result could well be an increase in racist attacks, ethnic conflicts, and an exodus of Gypsies to the “rich” West.

Prevention of this pessimistic scenario is not in the hands of health policymakers and medical researchers. Extensive educational programmes, control of the birth rate, new economic chances, and a change of lifestyle are the best that east European governments and Roma communities themselves could do for the health needs and the future of Roma people.

Emil Ginter  Head of Epidemiology
Institute of Preventive and Clinical Medicine, 85301 Bratislava, Slovakia


Citalopram is safe

Erratum—Power comments on my assertion that all selective serotonin reuptake inhibitors seem safe in overdose,4 citing six published cases of suicide in which citalopram was strongly suggested to have caused death. Although no drug is absolutely safe in overdose, the relative safety of the selective serotonin reuptake inhibitors compared with tricyclic antidepressants in this regard has been shown by many years of clinical experience.5

Citalopram is the most widely prescribed antidepressant in Sweden, and the Swedish Poisons Information Centre at the Karolinska Hospital has reported data on 104 cases of “pure” citalopram overdose treated in Swedish hospitals in 1995-6.6 Nausea, dizziness, and drowsiness were seen in patients who had ingested up to 30 times the usual therapeutic dose of 20 mg. In cases where up to 95 times the usual therapeutic dose was taken seizures and electrocardiographic abnormalities were reported, but there were no deaths or serious arrhythmias. The largest overdose in this series was equivalent to over 9 months’ treatment at 20 mg daily, showing considerable determination on the part of the patient, who made a full recovery.

The authors of the study concluded that most citalopram overdoses do not have an unfavourable course.7 An overdose of 4000 mg of apparently “pure” citalopram resulting in death8 was of similar size to a reported fatal overdose of fluoxetine taken alone.9 Power reasonably concludes that the true safety of a drug in overdose cannot fully be determined until the drug has seen extensive clinical use. Citalopram has been available in some countries since 1989, and it is estimated to have been used in routine clinical practice in almost 8 million patients. Thus there seems to me no reason to question the safety profile of citalopram any more than that of other selective serotonin uptake inhibitors.

A S Hale  Consultant psychiatrist
Thanet Community Mental Health Care, Westbrook Centre, Margate, Kent CT9 5DD

5 McKee M. The health of gypsies. BMJ 1997;315:1172-3. (8 November.)

Differences between Gypsies, Slovaks, and Hungarians living in Slovakia (district of Levice), 1996.

Figures are number (percentages) of positive answers

<table>
<thead>
<tr>
<th></th>
<th>Slovaks</th>
<th>Hungarians</th>
<th>Gypsies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (SD) (years)</td>
<td>37.9 (8.5)</td>
<td>38.4 (8.3)</td>
<td>38.0 (8.4)</td>
</tr>
<tr>
<td>Responses to questions asked</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Daily or almost daily consumption of:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Milk and milk products</td>
<td>286 (43.6)</td>
<td>140 (44.6)</td>
<td>12 (25.5)</td>
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<tr>
<td>Fruit</td>
<td>285 (43.5)</td>
<td>151 (48.0)</td>
<td>9 (19.2)</td>
</tr>
<tr>
<td>Vegetables</td>
<td>280 (42.8)</td>
<td>171 (54.5)</td>
<td>15 (31.9)</td>
</tr>
<tr>
<td>Beer</td>
<td>169 (26.8)</td>
<td>75 (23.9)</td>
<td>23 (49.0)</td>
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<tr>
<td>Spirits</td>
<td>67 (10.2)</td>
<td>39 (12.6)</td>
<td>11 (23.0)</td>
</tr>
<tr>
<td>Often in conflicts with other people</td>
<td>87 (13.3)</td>
<td>44 (13.8)</td>
<td>16 (34.0)</td>
</tr>
<tr>
<td>Agree with statement “I have no chance to influence my future”</td>
<td>73 (11.7)</td>
<td>34 (10.8)</td>
<td>13 (27.7)</td>
</tr>
<tr>
<td>Poor health status</td>
<td>51 (7.8)</td>
<td>22 (7.0)</td>
<td>10 (21.3)</td>
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<tr>
<td>Sleep disorders</td>
<td>53 (8.1)</td>
<td>30 (9.6)</td>
<td>8 (17.0)</td>
</tr>
<tr>
<td>Almost no reading of newspapers</td>
<td>178 (27.1)</td>
<td>55 (17.5)</td>
<td>23 (48.9)</td>
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<tr>
<td>No book read during past year</td>
<td>262 (40.0)</td>
<td>146 (46.5)</td>
<td>35 (74.5)</td>
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<td>Smoker</td>
<td>329 (50.2)</td>
<td>167 (53.2)</td>
<td>37 (78.7)</td>
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<tr>
<td>Father of more than two children</td>
<td>91 (13.9)</td>
<td>43 (13.7)</td>
<td>24 (51.1)</td>
</tr>
</tbody>
</table>

Removing intravenous lines at 72 hours allows need for antibiotics to be reassessed

Erratum—Randolph et al’s meta-analysis suggests that continuous infusions of low dose heparin prolong the patency of arterial and venous peripheral catheters.3 This is encouraging for both patients with needle phobia and junior doctors. The authors do not, however, emphasise the risks of invasive vascular devices, particularly the infective risks.

Although some studies disagree, it is widely accepted that rates of phlebitis and infection (both local and systemic) associated with intravenous lines are related to the duration of insertion.4 The American Hospital Infection Control Practices Advisory Committee strongly recommends...
replacing peripheral venous catheters in adults every 48–72 hours.

The problem of hospital acquired infection related to intravascular lines should not be underestimated. A recent survey by the Public Health Laboratory Service found that invasive devices constituted by far the most important risk factor for hospital acquired infection. From our own unpublished figures we found that 37% of cases of hospital acquired bacteraemia over two months were related to an intravascular line. Of these, 14% were associated with peripheral venous catheters; these catheters were probably underrepresented, because peripheral line tips are submitted for microbiological analysis infrequently.

In addition, the increasing rates of colonisation of hospital inpatients with resistant organisms, particularly methicillin resistant Staphylococcus aureus, are producing a concomitant increase in resistant bacteraemia related to lines. Analysis of bacteraemias within our trust over 17 months showed that 31% of all bacteraemias due to S aureus were resistant to methicillin.

Showing that the use of continuous infusions helps to decrease phlebitis is encouraging, and the decreased frequency of handling of the catheter hub site would reduce the risk of contamination—now a recognised factor in the pathogenesis of infection associated with intravascular devices. However, we would strongly advise against allowing the lack of visible inflammation to lead to continued use of peripheral intravenous lines. A policy of removing lines at 72 hours would allow consideration of the need for continued antibiotics or for switching to oral treatment with the attendant cost savings, patient comfort, and possible earlier discharge.

M Ruddy
Specialist registrar
C C Kibbler
Consultant
Department of Medical Microbiology, Royal Free Hospital, London NW3 2QG.

1 Randolph A, Cook DJ, Gonzales CA, Andrew M. Benefit of handling of the catheter hub site would reduce the risk of contamination—now a recognised factor in the pathogenesis of infection associated with intravascular devices. However, we would strongly advise against allowing the lack of visible inflammation to lead to continued use of peripheral intravenous lines. A policy of removing lines at 72 hours would allow consideration of the need for continued antibiotics or for switching to oral treatment with the attendant cost savings, patient comfort, and possible earlier discharge.

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M Ruddy
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C C Kibbler
Consultant
Department of Medical Microbiology, Royal Free Hospital, London NW3 2QG.
CAGE questionnaire allows doctors to avoid focusing on specifics of drinking

Editor—Perhaps I should be flattered that my CAGE questionnaire is so popular that the BMJ printed it twice in 1997—though without proper attribution on both occasions. One of the articles by Ashworth and Gerada in the ABC of Mental Health uses the CAGE questionnaire as if it had just appeared out of nowhere rather than being the result of clinical studies and efforts to develop a brief and easily remembered series of questions.1 The citation should have been given.2 Ashworth and Gerada should read a paper by Steinweg and Worth, who write: “Our study demonstrates the negative impact of asking patients to discuss the specifics of their drinking. Not only can it make for an uncomfortable interview, it is diagnostically self-defeating. The CAGE questions offer the physician powerful tools to avoid focusing on the specifics of drinking. However, important keys to the CAGE are open-ended introductions and resisting the urge to ask the patient to quantitate consumption.” Ashworth and Gerada do the opposite in their recommendations. After a focus on amount, time, and pattern of consumption they say: “Specific questioning should continue with the CAGE questionnaire.” I think that study of Steinweg and Worth’s paper will persuade them to change their recommendations.

John A Ewing
Professor emeritus of psychiatry, University of North Carolina
2311 Canterwood Drive, Wilmington, NC 28401, USA


Audit Commission tackles anaesthetic services

Anaesthesia should remain physician based service.

Editor—We read with interest the articles1 2 that you recently published in response to the Audit Commission’s Anaesthesia under Examination,3 which revisits the topic of nurse anaesthetists. We trained as anaesthetists working both in and out of operating theatres time to concentrate on those patients who require the benefit of their medical training and extended specialist experience.

D N Robinson
Fellow in paediatric anaesthesia
HGC (Scotland), Clydebank G81 4HX

1 Smith A. Audit Commission tackles anaesthetic services. BMJ 1998;316:3-4. (3 January.)

Quoted paper did not say that participation of nurses makes adverse outcomes more common

Editor—In his editorial1 Smith quoted a paper that I cowrote2 to support his argument that adverse outcomes in anaesthesia are more common when anaesthetists are given by nurse anaesthetists.

Our article did not mention anything of this sort. We had conducted a retrospective analysis of all reports that the department of a university hospital made to the faults, accidents and near accidents committee of that hospital over 10 years.

D N Robinson
Fellow in paediatric anaesthesia
HGC (Scotland), Clydebank G81 4HX

1 Smith A. Audit Commission tackles anaesthetic services. BMJ 1998;316:3-4. (3 January.)

Non-physician anaesthetists may free up consultants to concentrate on patients requiring special attention

Editor—Smith’s editorial gives an overview of the Audit Commission’s report on anaesthetic services.1 Inevitably, however, headline reporting in both the lay and the medical press concentrated on the report’s recommendation to consider the introduction of non-physician anaesthetists into British practice. Smith points out the firm opposition of the Association of Anaesthetists to this—a result of concerns over the safety of patients, legal ambiguities, and cost.2 Rather than being seen as a threat, however, the report should act as an impetus for anaesthesia in the United Kingdom to reconsider this entrenched, “closed shop” attitude.

Much routine minor and intermediate surgery performed on fit patients does not require the presence of a fully trained consultant anaesthetist for the entire duration. The training guidelines of the Royal College of Anaesthetists suggest that, after an introductory module of 12 weeks, medically qualified trainees may have “level 2” supervision for some straightforward cases. This is defined as: “Trainer present in the operating theatre suite … available to assist or advise.”

Historically, much routine work has been done by trainee anaesthetists under the (sometimes fairly distant) supervision of consultants. As a result of the Calman reforms and limits on working hours this service contribution from trainee doctors is being reduced drastically. At the same time, the demands on anaesthesia services are growing constantly. The answer to this problem should not be simply to seek ever increasing numbers of consultant anaesthetists to perform undemanding work, but rather to encourage the development of non-physician anaesthesia practitioners, trained and supervised by medically qualified anaesthetists. Surely, a nurse or other medical professional who would undertake two or three years’ training in anaesthesia could provide care at least equivalent to that of a trainee doctor.

This would provide consultant anaesthetists working both in and out of operating theatres time to concentrate on those patients who require the benefit of their medical training and extended specialist experience.
We did not analyse the data on the basis of who gave the anaesthetics. This was irrelevant for our study since it is the practice in all university hospitals in the Netherlands that anaesthesia is given by a team consisting of an anaesthesia trainee and an anaesthetic nurse under the direct supervision of a qualified specialist anaesthetist. It is unfortunate that our study has been misquoted in support of the argument that nurse anaesthetists are not as safe as medical anaesthesiologists.

V Chopra  Consultant anaesthetist
Leiden University Medical Centre, 2300 RC Leiden, Netherlands

1 Smith A. Audit commission tackles anesthetic services. BMJ 1998;316:3-4. (3 January)

Investment is required to increase number of consultant anaesthetists

Editor—I am used to seeing misrepresentative headlines in the tabloid press but am surprised to see one in the BMJ.1 The Audit Commission’s wide ranging report Anaesthesisia under Examination makes no mention of urging the NHS to appoint nurse anaesthetists as the title of Wise’s article suggests.2

The report does recognise that one obvious way to reduce the demand for more doctors substantially is to allow non-medical staff to administer anaesthesia. This cannot be argued with. The actual recommendations, however, are much more tentative and suggest that hospitals such as small district general hospitals, with few trainees or perhaps none at all, might benefit from a trial of using non-physician anaesthetists (or physicians’ assistants, as they are called in some parts of the United States) to help the lone consultant. This would involve an extension of the role of the anaesthetic nurse or operating department assistant to a more active role in the maintenance of anaesthesia. In certain circumstances this might lead to the consultant supervising two operating theatres at one time, with suitable staff and a suitable case mix.

This system might well be acceptable to some consultants without any trainees, who must be willing to take on the additional responsibility and should be paid appropriately to do so. But it is patently not the traditional role of the nurse anaesthetist in the United States, where many small hospitals are virtually autonomous and only nominally under the charge of a surgeon, let alone an anaesthetist. I suspect that few patients in the United Kingdom would be willing to accept this arrangement.

A survey of staffing in anaesthesia showed that the United Kingdom (and the Republic of Ireland) has by far the lowest number of anaesthetists in the UK, compared with 10.8 in 17 European countries.3 Despite the much higher number of consultants (with few trainees) in the rest of Europe, however, greater use is made there of anaesthesia nurses (15.5/100 000 in Germany and about 23/100 000 in Norway and Sweden). So no cost savings are made.

Thus international evidence shows clearly that as (or if) the number of trainees is reduced a massive investment is required in consultant anaesthetists to bring the United Kingdom into line with the rest of Europe. Only when this has occurred is a trial with non-physician anaesthetists warranted.

D W Green  Consultant anaesthetist
King’s College Hospital NHS Trust, London SE5 9RS

1 Wise J. NHS urged to appoint nurse anaesthetists. BMJ 1998;316:10. (3 January)

Experience is paramount for successful management of disease

Editor—Earlier this week the medical firm on which I am registrar was on general medical intake. After a quiet day things began to get busy around 11 pm. Two patients with suspected oesophageal varices and haematemesis arrived almost simultaneously. Both had signs of cardiovascular compromise. Around midnight a woman with an anterior myocardial infarct was given thrombolysis. At 2 am the same patient developed transient complete heart block and pulmonary oedema, then ventricular tachycardia followed by atrial fibrillation. Too short of breath to lie flat for transvenous pacing, she was successfully managed with an external pacing device, followed by intravenous amiodarone and carefully titrated doses of intravenous nitrates.

Our firm comprises one house officer, two senior house officers, and a registrar, and at the time of writing it had a third senior house officer as well. We managed all of the ill patients described above. The duty senior house officer was clerking patients who had taken overdoses or had other less acute problems for most of the night and was therefore unable to be present with me. As a fairly experienced registrar I have managed similar cases countless times before. I am experienced at staying awake and managing patients “hands on” when needed, as was no doubt the case here.

What is disturbing is that not one of the four doctors junior to me on the firm learnt anything about the management of patients with acute serious illness that night. In three years from now I might be a consultant. Those doctors, deprived of experience by the new deal, may be medical registrars. My own mother will be approaching the same age as the patient with the infarct. From the time of Hippocrates experience has been recognised as paramount for the successful management of disease. Are our mothers and fathers to be managed by inexperienced doctors?

Martin Dumskij  Specialist registrar in general medicine, diabetes, and endocrinology
Kingston Hospital, Kingston upon Thames KT2 7QB

Urgency and priority models

Model has limited practical application

Editor—The recent interest in priority scoring for surgical operations is to be commended. The paper by the Northern Ireland Clinical Resource Efficiency Support Team highlights the differences between “urgency” and “priority” and how demographic and lifestyle related details may influence the timing of coronary artery bypass surgery.4 Unfortunately this model has limited practical application.

Firstly, although the opinion of the referring doctor is important, the ultimate responsibility for prioritising the waiting list lies with the cardiac surgeons, who represented only a minority of the judges (4 out of 33). Furthermore, the study does not take into account how cardiac surgeons organise their waiting lists. Some run their list individually and some operate a joint waiting list; most would separate the list into emergency, urgent, and routine cases, but few would adopt a more sophisticated priority system within each group. This is not to say such an approach would be wrong, but it would imply a constant reshuffling of patients as the list grew and would make it impossible to give patients a reasonable estimate of their wait.

Secondly, inspection of the angiogram is an integral part of patient assessment, but this was denied to the judges in this study. Simply recording the number of severely diseased vessels fails to take into account the importance of the anatomical location of the stenoses, the dominance of the coronary vessels, and their relative size. A stenosis of the proximal left anterior descending artery in association with a tight proximal circumflex stenosis would be assigned more urgency and priority than perhaps a stenosis of the distal left anterior descending artery—in combination with disease in the posterior descending artery—yet both are examples of two vessel disease. This point is particularly identified in a study from New Zealand.5

Finally, the success of interventional cardiology creates an increasing number of grey areas. The option to treat surgically or by endovascular intervention may be biased by the relative length of the waiting list for each technique and by the patient’s own wishes, as much as by the clinical need for intervention. Although the model presented allows those deciding about waiting lists some insights into how others view the surgical treatment of coronary artery disease, the decisions made were based on insufficient clinical data and may not reflect current practice. Unfortunately, this model also fails...
to recognise the key role of the cardiac surgeon as the “gate keeper” of the waiting list.

Ian M Mitchell  Consultant cardiothoracic surgeon
David W Quinn  Research fellow
Nottingham City Hospital, Nottingham NG5 1PB


Distinction between urgency and priority helps no one

Entror—Kee et al distinguish between urgency and priority in patients waiting for coronary artery bypass surgery.2

In practice such a distinction is so fine as to be pointless; the most urgent patients receive the highest priority. It is not necessary to convene workshops and invoke paper patients to determine urgency. The amount of resources on coronary artery bypass surgery allows the invocation of survival benefits over medical treatment to be defined precisely.3

For example, patients with three vessel coronary artery disease and angina of class III or IV have a one year survival of 94% after coronary artery bypass surgery and of 85% if treated medically.2 If these patients are kept on a waiting list for one year twice as many women aged over 64 will have died than those operated on for symptoms alone.4 Nevertheless the 25 patients with three vessel coronary disease and angina of class III or IV angina still waited a median of 81 days (interquartile range 45-127) to be operated on.5

We have analysed a random sample of 67 patients from one health district who underwent coronary artery bypass grafting between 1 April 1996 and 31 December 1997. The reason for the operation was to improve prognosis, regardless of the severity of symptoms (class I), or to improve symptoms alone, as defined by the American College of Cardiology and American Heart Association (class II).6 In 56 patients the bypass operation conferred a survival advantage. These patients had a shorter wait than those operated on for symptoms alone (table). Those with the most threatening coronary artery disease and the worst symptoms had the shortest wait (table). Nevertheless the 25 patients with three vessel disease and class III or IV angina still waited a median of 81 days (interquartile range 38 to 169).

We believe that these data show that the priority given to these patients is appropriate, given what is known of the clinical course of coronary artery disease. We also believe that some of these waiting times are unacceptable. No amount of “clinical judgment analysis” can alter this. Waiting lists can be reduced only by increasing the number of operations or denying surgery to certain patients who currently benefit. Muddying the waters with academic distinctions between urgency and priority and the effect that “lifestyle characteristics” have on this helps no one—our patients least of all.

Philip Wrage‡  Medical student
University of Sheffield, Sheffield S10 2RX

Walter Rhodes  Consultant cardiologist
Barnsley District General Hospital, Barnsley S75 2EP

Graham Cooper  Consultant cardiothoracic surgeon
Northern General Hospital, Sheffield S5 7AJ


Clinical and economic arguments favour extension to upper age limit for breast screening

Entror—In an editorial Sutton argues that the upper age limit of 64 for breast screening is illogical, and he ascribes it to ageism.7 This may be true, but there is no economic rationale for limiting routine screening to women below the age of 65. Indeed, the most cost effective age in terms of cost per life year saved by breast screening is 70.8 We have recently set out the economic arguments for extending breast screening to women aged over 64; some of these have been repeated. Firstly, there is the issue of poor compliance with screening. The level of compliance with any screening programme is an economic issue only if those who do not comply are at a higher risk than those who do, which for breast screening does not seem to be the case.9 Secondly, poor compliance is sometimes wrongly associated with non-attendance. Non-attendance wastes screening resources, but if the national breast screening service were to use an efficient appointment method that required women to confirm their intention to attend then non-attendance could be reduced to negligible levels.10 Thus absolute levels of compliance for breast screening are largely irrelevant for cost effective screening.11 For instance, if compliance of 90% produces a fall in deaths from breast cancer of 30% then a compliance level of 45% that reduces deaths by 15% at half the cost is still cost effective.12 The reason that breast screening has not been extended to older women is probably confusion of screening objectives rather than ageism. Compliance with screening has been implicitly and explicitly set as a key objective when really the ultimate objective of screening should be a reduction in mortality and morbidity from the disease.13 One reason why screening of older women has not been adopted, despite the overwhelming clinical and economic arguments for it, is resource limitation. This, however, can be addressed by simply abandoning screening for women aged 50-53 and redirecting the resources to women aged 66-69. Such a policy would increase the numbers of life years saved by 5% at no extra cost.14

David J Torgereson  Senior research fellow
National Primary Care Research and Development Centre, Centre for Health Economics, University of York, York YO1 5DD

Toby Gosden  Research fellow
National Primary Care Research and Development Centre, University of Manchester, Manchester M13 9PL

1 Sutton GC. Will you still need me, will you still screen me, when I’m past 64? BMJ 1987;315:1065-6. (25 October.)

Gamete donors for IVF should relinquish right of ownership to resulting embryos

Entror—The Human Fertilisation and Embryology Act states that gamete donors have rights over the destiny of their gametes and any embryos resulting from their use.1 Donors are volunteers, who are given a full explanation of the use of their gametes for clinical or research purposes. After counselling, donors proceed to donation only if they are in full agreement about the use of their gametes, which extends to deciding the fate of any resulting embryos from their donation.

The length of time that embryos can be stored by cryopreservation has recently been extended from five to 10 years, provided that the natural parents of the frozen embryos consent to this. It is essential that both parents agree about the length of cryopreservation. In cases in which one of the natural parents is a donor, his or her agreement must be obtained before the fate of the frozen embryos is decided, whatever the other natural parent may wish.

We recently encountered a difficult situation. A couple receiving in vitro fertilisation with donated sperm had eight...
embryos frozen at the end of treatment, which were still frozen at the end of the initial five year storage period. The donor could not be contacted to provide consent to further storage. Although the female partner was the natural mother of the frozen embryos, because the sperm donor could not be contacted and the patients did not wish to put the matter before the law an extension to embryo storage could not be allowed and the embryos were destroyed.

To prevent this situation arising again, we believe that after treatment the responsibility for donated gametes should be accepted by the patient or couple to whom they are donated. The fate of embryos resulting from gamete donation should be decided solely by the patient or couple, after counselling.

A change in the Human Fertilisation and Embryology Act is necessary to reflect these views and to strengthen the rights of patients as to the fate of donated gametes and resulting embryos. Gamete donors consenting to the use of their gametes for clinical purposes should relinquish the right of ownership for any resulting embryos to the couple for whom the embryos are intended. This will effectively give the patients sole authority over the fate of frozen embryos.

M R Gazvani Research fellow
A J M Thomson Research fellow
S J Wood Research fellow
C R Kingsland Director, consultant obstetrician and gynaecologist
Reproductive Medicine Unit, Liverpool Women's Hospital, Liverpool L8 7SS
D L Lewis-Jones Senior lecturer in obstetrics and gynaecology
Department of Obstetrics and Gynaecology, University of Liverpool, Liverpool


Water fluoridation is safe and effective

Editor—It was not surprising that Gibson and Gibson1 had to rely on a paper published 50 years ago to support their claim that water fluoridation was not effective in preventing tooth decay and that it was harmful to health. However, the paper they cited2 did not mention either a delay in tooth eruption or the occurrence of pulmonary embolism, which is a clear indication for insertion of a filter. The indications commonly treated in this way in the United Kingdom are proven pulmonary embolism occurring despite adequate treatment with anticoagulant drugs, contraindications to treatment with anticoagulant drugs with demonstrable thrombosis in the femoral or iliac veins, complications of treatment with anticoagulant drugs requiring discontinuation of treatment, or proximal thrombosis in the iliac vein or vena cava in patients about to undergo surgery (for example, pelvic reconstruction after trauma).2

While it is true that randomised studies are lacking, it is equally true that it would be difficult ethically to assign patients randomly to treatment given the accepted indications in the United Kingdom. The study shows that filters reduce but do not eliminate the occurrence of pulmonary embolism, which is encouraging but it did not find a reduction in mortality. This would suggest that filters should not be employed as first line treatment in the population that is studied; this finding is not surprising. I do not believe, however, that this data should be used to discourage the use of filters to treat those indications described above since the study design did not address those issues.

Mark G Cowling Lecturer in interventional radiology
Division of Radiological Sciences, United Medical and Dental Schools of Guy's and St Thomas's Hospitals, Guy's Hospital, London SE1 9RT

1 Minerva. BMJ 1998;316:788. (7 March.)

Providing letters to patients

Patients find summary letters useful

Editor—Essex raises some interesting points about the value of giving patients a written summary of their consultation.1 The practice of sending patients a letter summarising their consultation is very common in genetic counselling. A colleague and I have completed a qualitative interview study exploring patients’ attitudes about and their use of written summaries of their genetic consultations for hereditary breast and ovarian cancer.2 Like Essex, we found that patients responded very positively to these letters. Altogether 37 (95%) out of 40 patients in the sample said that the summary letter aided their understanding or recall of information that had been given in the clinic, or both. The written summary was also perceived as valuable because it could be shown to other clinicians to support the patient’s case for gaining access to breast or ovarian screening programmes, it reassured patients that they were taking appropriate action, and it contained information about other relatives’ risks. In addition, the written summary was also perceived as a useful tool for disseminating genetic information to other family members; 34 (85%) out of 40 patients said they had used, or intended to use, the written summary of their counselling session to facilitate the communication of genetic information to other biological relatives.

On the basis of our findings we suggest that genetic counsellors send patients a letter summarising their consultations as this may result not only in an increase in the patient’s understanding, but may prevent the miscommunication of genetic information within the family. However, we feel that clinicians should be aware that providing patients with a written summary of their consultation may also have negative consequences. It may generate an inappropriate demand for referrals to genetic clinics from family members at low risk, and, more importantly, reading a letter written to the patient may cause needless anxiety among low risk family members.

N Hallowell Senior research associate
Centre for Family Research, Faculty of Social and Political Sciences, University of Cambridge, Cambridge CB2 1RF

1 Essex C. Consultants could give patients a letter summarising their consultation. BMJ 1998;316:786. (28 February.)

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Letters
Letters

GPs can be given copies of letters sent to patients

Editor—I would like to reinforce Essex’s opinion that consultants should write letters to their patients summarising their findings after a consultation.1 Since starting the breast screening service in South Essex in 1990, it has been my policy to write to women who have been assessed and whose abnormalities have been found not to require surgical evaluation or treatment.

Initially, I also wrote letters to the woman’s general practitioner but I have since found this to be an unnecessary waste of resources. I now write to the woman and send a copy of the letter to her general practitioner. This ensures that the general practitioner knows exactly what the woman has been told and it provides a record of the procedures undertaken and the advice given on future management.

There has been a mixed reception to the letters. One group of general practitioners praised their clarity while another complained about their technical detail. I hope modifications made in response to these comments have made the letters more comprehensible to both the women and their general practitioners.

M D Lewars Consultant radiologist
South Essex Breast Screening Service, Westcliff on Sea, Essex SS0 0SB

1 Essex C. Consultants could give patients a letter summarising their consultation. BMJ 1998;316:706. (28 February.)

Summary letters may be especially appropriate after emergency admissions

Editor—It is common for professionals such as accountants and solicitors to follow a meeting with a client with a letter recapitulating the points discussed, and Essex is to be congratulated on his initiative in doing the same after paediatric outpatient appointments.1 Communication is especially important at either end of the age spectrum, and some years ago two of the four geriatric teams in our hospital decided to send out letters to patients who had been seen for consultations. The letters were sent after the initial visit and also when results came back or decisions were arrived at which might have made a further letter helpful. Altogether, 75% of 35 patients found the letters “very helpful” and 18% found them “helpful.” Questionnaires were also sent to 38 patients in a control group. In this group, 74% indicated that a letter would have been “very helpful” and 23% indicated that it would have been “helpful.” We sent separate letters to the patients’ general practitioners together with a copy of the letter sent to the patient.

It was a valuable exercise in communication skills, but we had to discontinue the practice due to the need to reduce the secretarial (and medical) workload. Perhaps it would be even more useful to send a similar letter after an emergency admission, since so much of what has happened becomes eclipsed by subsequent events, and memories of procedures and discussions so readily become blurred.

Nicholas Coni Consultant geriatracian
Department of Medicine for the Elderly, Addenbrooke’s Hospital, Cambridge CB2 2QQ

1 Essex C. Consultants could give patients a letter summarising their consultation. BMJ 1998;316:706. (28 February.)

Advice to authors

We prefer to receive all responses electronically, sent either directly to our website or to the editorial office as email or on a disk. Processing your letter will be delayed unless it arrives in an electronic form.

We are now posting all direct submissions to our website within 72 hours of receipt and our intention is to post all other electronic submissions there as well. All responses will be eligible for publication in the paper journal.

Responses should be under 400 words and relate to articles published in the preceding month. They should include ≤5 references, in the Vancouver style, including one to the BMJ article to which they relate. We welcome illustrations.

Please supply each author’s current appointment and full address, and a phone or fax number or email address for the corresponding author. We ask authors to declare any conflicts of interest.

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Cover picture meant that BMJ had descended to level of tabloid newspapers

Editor—It was with deep regret that I saw the cover picture for the issue of 6 June. The issues raised by the Bristol case are matters that have grave implications for the whole of the profession.1 It raised many difficult questions, which the BMA and other professional bodies are seeking to address both in depth and with great urgency.

To sensationalise the issue is unnecessary and deeply offensive to many members of the association. Consultants are only too aware of the great personal tragedy that has affected the families of the children in the Bristol case but nevertheless will be saddened to see a learned journal descend to the level of the tabloid newspapers.

J N Johnson Chairman, Central Consultants and Specialists Committee
BMA, London WC1H 9JP


2 The BMJ publishes cover pictures to draw attention to the journal publishing a cluster of material on a subject. Our experience is that readers like the occasional use of cover pictures. When we do publish a cover picture we find as strong an image as we can to illustrate the material. Last week’s issue contained nine papers that moved on the debate that the Bristol case has started, a debate that may prove to be one of the most important in British medicine this century. We wanted to ensure that readers knew about these articles, and so we decided to publish a cover picture. It was an obvious thing to do to use a strong picture associated with the Bristol case.

Whether or not the cover is tabloid is a matter of opinion; to my mind it simply shows a grieving mother. But nobody could argue that the content, which is what matters most, is tabloid. And the good thing about tabloids is that millions of people read them, whereas the bad thing about many scientific journals is that nobody reads them, not even scientists—Editor.

J A Eaden Research fellow
B Ward Audit assistant

J F Mayberry Consultant physician
Gastrointestinal Research Unit, Leicester General Hospital, Leicester LE1 4PW

1 Essex C. Consultants could give patients a letter summarising their consultation. BMJ 1998;316:706. (28 February.)

