

children to receive intensive care in lead centres^{1,2} and this might be expected to reduce mortality across the board.

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References

- 1 Pearson GA, Shann F, Barry P, *et al*. Should intensive care be centralised? Trent versus Victoria. *Lancet* 1997;**349**:1213–17.
- 2 Pearson G, Barry P, Timmins C, *et al*. Changes in the profile of paediatric intensive care associated with centralisation. *Intensive Care Med* 2001;**27**:1670–3.

Improved outcome in severe meningococcal disease

We thank Tibby *et al* and Pearson for their interest in our paper.¹ We agree with Pearson that evidence for a fall in overall mortality in meningococcal septic shock would require a geographical community based study. We described mortality in severe meningococcal disease in a paediatric intensive care unit (PICU).

In our multispeciality PICU in the north west, we have observed a continued decrease in both actual PICU mortality and mortality adjusted for disease severity since the original study period (table 1). Paediatric index of mortality (PIM) is a more contemporary scoring system than PRISM (paediatric risk of mortality score), and so has been calibrated to the more recent decline in PICU mortality rates.² PIM gives a score at point of first PICU contact.

This general trend of improving meningococcal outcome is also reflected in other PICUs. As shown by the results from St Mary's PICU in London, where in a group with an overall actual mortality of 18.7% (PICU mortality for the study period being 10%, and an additional 8.7% mortality for the "unretrievables"), they encouragingly had managed to reduce the meningococcal PICU mortality in their "specialist PICU" from 23% to 2% (1992–97).³ Tibby *et al*, from Guy's Hospital PICU in London (1998–2001), in their letter report a similar very low mortality rate.

There has been continued improvement in outcome from severe meningococcal disease throughout the UK. Early recognition and early institution of treatment are of paramount importance. No single centre holds the monopoly on the improved outcome in meningococcal disease. Although improved intensive care has undoubtedly contributed to this fall in mortality, there should be more recognition of the role of those in the community, parents and carers, general practitioners, and district general hospitals who have significantly contributed (and continue to contribute) to the survival of these critically ill children.

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Table 1 Actual mortality, number of patients, mortality per year, and standardised mortality ratio (SMR) in patients admitted to the paediatric intensive care unit (PICU) at the Royal Liverpool Children's Hospital with meningococcal disease

| | Jan 1995 to March 1998 ¹ | April 1999 to March 2001 |
|----------------------------------|-------------------------------------|--------------------------|
| Actual mortality/PICU admissions | 11/123 (8.9%) | 3/95 (3%) |
| Mortality per year | 3.5 | 1.5 |
| PIM predicted SMR | 1.16 | 0.24 |

SMR = paediatric index of mortality (PIM) predicted mortality/actual mortality.

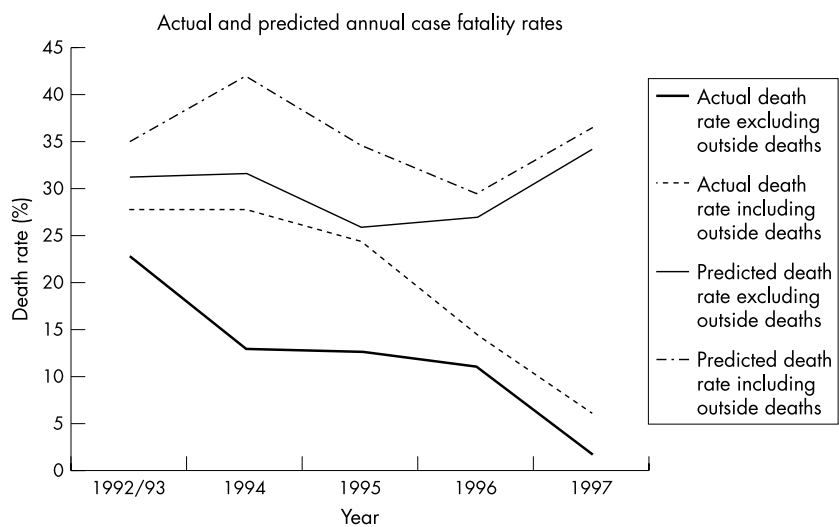


Figure Actual and predicted annual case fatality rates.

References

- 1 Thorburn K, Baines P, Thomson A, *et al*. Mortality in severe meningococcal disease. *Arch Dis Child* 2001;**85**:382–5.
- 2 Shann F, Pearson G, Slater A, *et al*. Paediatric index of mortality (PIM): a mortality prediction model for children in intensive care. *Intensive Care Med* 1997;**23**:201–7.
- 3 Booy R, Habibi P, Nadel S, *et al*. Reduction in case fatality from meningococcal disease associated with improved healthcare delivery. *Arch Dis Child* 2001;**85**:386–90.

Mortality in meningococcal disease: please report the figures accurately

We thank Tibby and colleagues for their interest. We believe they and others would be interested in the accompanying figure (see above).

It compares yearly case fatality rates on all referrals to St Mary's PICU, regardless of whether they died before a mobile intensive care team arrived or while the team was assisting with resuscitation. The 29 "outside" deaths are included (3 in 1992/3, 8 in 1994, 10 in 1995, 3 in 1996, 5 in 1997). As stated in our published paper, logistic regression analysis, controlling for disease severity, age and sex, and including these extra deaths, showed no change in the estimated odds ratio for the yearly reduction in death rate, namely 0.41. The overall case fatality rate for 1997 became 6% compared with the PICU admission rate of 2% and a predicted case fatality rate 34% using PRISM scores.

For the 5 deaths in 1997 outside St Mary's PICU, response times between call to the unit and arrival of a team at the DGH varied between 100 and 185 minutes. One child died

as the local hospital were telephoning us, two arrested within 90 minutes of St Mary's being called and died within minutes of the team arriving, and the other two died between 2 and 7 hours after arrival.

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Genuine reduction in meningococcal deaths results from teamwork

As paediatric intensivists in lead centres accredited for paediatric intensive care (PIC) training and responsible for the care of approximately 7000 cases per year, we read with concern the report from St Mary's Hospital which reported improved outcome of meningococcal disease (MD) in 1997 compared with previous years.¹

Their reported reduction in mortality must be seen in the context of an overall reduction of childhood mortality and a widespread improvement in the outcome for many conditions requiring PIC such as acute respiratory failure,² persistent pulmonary hypertension³ and complex congenital heart defects.⁴ Overall UK PIC mortality rates have fallen to a standardised mortality ratio (SMR) of 0.87 as assessed by the Paediatric Index of Mortality⁵ compared with the model generated in 1994.⁶

Their application of the severity of illness score (PRISM) is incorrect. No patient has a 100% predicted risk of mortality and therefore all deaths observed in any such study must increase the SMR. The exclusion of nearly half of the total deaths (29/62, 47%) who did not survive the long stabilisation and overall retrieval times must reduce SMR regardless of any other intervention. Whilst inclusion of these cases does not alter the direction of the relationship between SMR and year, it raises the overall mortality in the series towards 20% and more than doubles the headline mortality in 1997. Data from the last 4 years would be of interest. In addition, the lack of any data relating to the performance of the model in different risk groups fails to address the potential confounding factor of disease severity. Since all survivors will reduce SMR, one cause of apparent improvement in risk-adjusted survival is increased admission of low risk cases.

Recent series from other institutions have followed the convention of presenting data by level of predicted risk.⁷⁻⁹

The claim that their "Mobile Intensive Care" service is the key element in improved survival is confusing when all the cases that died under the care of this service were excluded from both the analysis and the "headline" figure of 2% mortality for MD.

However, our greatest concern is the claim that these data support their particular "model" of care of critically ill children. This is not consistent with their report, as St Mary's has not been performing transports since 1992 but the fall in mortality occurred some 4-5 years later. It should be remembered that PICU retrievals have been performed in Liverpool and Glasgow since the late 1970s. Their claim that this "model" has reduced mortality of meningococcal disease is also inconsistent with the similar improvements in outcome presented by other PICUs.⁷⁻⁹

We feel the narrow focus of the paper on the ICU care of MD is misleading. It ignores the important contribution of many others including parents, charities, and healthcare workers. Their role in education, early identification, treatment, and immediate high quality resuscitation is discounted. To imply that ICU management after the initial resuscitation is the key factor in improved survival undermines the vital contributions of these groups.

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References

- Booy R, Habibi P, Nadel S, et al. Reduction in case fatality rate from meningococcal disease associated with improved healthcare delivery. *Arch Dis Child* 2001;**85**:386-90.
- Peters MJ, Tasker RC, Kiff KM, et al. Acute hypoxemic respiratory failure in children: case mix and the utility of respiratory severity indices. *Intensive Care Med* 1998;**24**:699-705.
- UK collaborative randomised trial of neonatal extracorporeal membrane oxygenation. UK Collaborative ECMO Trail Group. *Lancet* 1996;**348**:75-82.
- Andrews R, Tulloh R, Sharland G, et al. Outcome of staged reconstructive surgery for hypoplastic left heart syndrome following antenatal diagnosis. *Arch Dis Child* 2001;**85**:474-7.

- Pearson GA, Stickley J, Shann F. Calibration of the paediatric index of mortality in UK paediatric intensive care units. *Arch Dis Child* 2001;**84**:125-8.
- Shann F, Pearson G, Slater A, et al. Paediatric index of mortality (PIM): a mortality prediction model for children in intensive care. *Intensive Care Med* 1997;**23**:201-7.
- Thorburn K, Baines P, Thomson A, et al. Mortality in severe meningococcal disease. *Arch Dis Child* 2001;**85**:382-5.
- Peters MJ, Ross-Russell RI, White D, et al. Early severe neutropenia and thrombocytopenia identifies the highest risk cases of acute meningococcal disease. *Ped Crit Care Med* 2001;**2**:225-31.
- Tibby, S. M. Mortality in meningococcal disease: please report the figures accurately. <http://adc.bmjournals.com/cgi/eletters/archdischild;85/5/386#224>. 26-10-2001.

Reduction in case fatality rate from meningococcal disease is due to genuine teamwork

We read with disappointment the response of Dr Peters and colleagues¹ to our article "Reduction in case fatality rate from meningococcal disease associated with improved healthcare delivery".² It is unfortunate that there appears to be a misunderstanding of the message of our study which demonstrated a significant improvement in the mortality of children with meningococcal disease (MD) over a period of time. Contrary to their concerns those results were achieved through genuine teamwork³ as stated in our paper.

In answer to the specific points they raised: We and other intensivists are also aware that mortality in conditions other than MD is also improving. In our paper we did not state that MD was the only condition in which there is an improvement in mortality. Our paper referred to a study published in *Critical Care Medicine* which also showed improving survival rates of paediatric patients (with various diseases) over time in another paediatric intensive care (PIC) setting.⁴

With reference to the patients who died at the referring hospital and their exclusion from the study. Our paper clearly states "Logistic regression analysis, controlling for disease severity, age and sex, showed that over the study period (1992-97) the overall estimate for the reduction in the odds of death was 59% per year (odds ratio for the yearly trend 0.41, 95% CI p=0.00001). This estimate and significance remained the same after inclusion of the 29 deaths that occurred at local hospitals".

We did not claim that mobile intensive care is the key element in improved survival. What we stated was: "Considerable changes in the management of patients with MD have occurred over the study period. While no single factor alone is likely to explain the reduction in mortality, several factors might have contributed to the improved outcome. In the past, few centres, including those with PICUs, admitted more than a small number of patients with MD annually. Furthermore, patients were often considered too sick to transfer to a specialist centre and were treated in the A&E department, paediatric ward or adult ICU of the local district general hospital. Establishment of a mobile intensive care team allowed the centralisation of care of children with MD at a specialist clinical and research unit, which in turn enabled extensive experience in the management of MD to be developed; this may be the most important reason for the improved outcome.... In other words, it was the increased experience in

dealing with meningococcal disease that was the critical factor.

The role of mobile intensive care was more directly addressed when we stated that it "has probably been another important factor in improved outcome", not the key factor.

The conclusions of our paper clearly state the multiple factors responsible for the results of the study, which have shown that a notable reduction in the case fatality rate for MD has been achieved.

The purpose of presenting our data was to emphasise the improvements in mortality in a particular group of patients brought about by a change in health care delivery. The key point being early intervention by a multidisciplinary team with a major research interest in the care of the critically ill child with infectious disease, who have the benefit of a "critical mass" experience.

The PICU at St Mary's Hospital, London was established in 1992, at the time primarily to facilitate the enrolment of children with meningococcal disease into clinical trials. As a large number of critically ill children were referred to our unit, we were subsequently able to record high-quality data regarding clinical status, severity of illness and outcome. We began to demonstrate a reduction in mortality from 1994 onwards, as it takes time to establish the clinical experience which can have a significant impact on the disease process.

The unit at St Mary's has been greatly involved in the development of a model of care involving "genuine teamwork" with the aim of improving the healthcare of children with MD. To this end we have been working with the meningitis charities (which are acknowledged on the paper) and other agencies to develop guidelines, publish treatment algorithms and improve policies. In addition our research unit has played a key role in the design and implementation of clinical trials of adjunctive treatments in meningococcal disease, which has led to the publication of the only two large randomised, double blind, placebo controlled studies in childhood septic shock.^{4,5}

Finally we are humbled by the magnitude of response from many other colleagues who have applauded our efforts. We believe, and have repeatedly stated, that what has been widely accepted as a major advance in the outcome of children with MD, could only have been achieved by multidisciplinary effort involving all sectors of health care delivery.

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References

- Petros AJ, Weir P, Marsh M, et al. Genuine reduction in meningococcal deaths results from teamwork. [electronic response to Booy et al, Reduction in case fatality rate from meningococcal disease associated with improved healthcare delivery]. www.archdischild.com <http://adc.bmjournals.com/cgi/eletters/archdischild;85/5/386> (accessed 7 March 2002).