Severe rombergism due to gentamicin toxicity

Sir,—Drs Roderick Duncan and Ian D Melville (31 October, p 1141) describe a case of rombergism with hearing loss in a patient who was an elderly man who received gentamicin 80 mg three times a day intramuscularly for eight days and a second similar course lasting six days. The serum gentamicin concentration of 13-6 mg/l during the second course was presumably a trough value. No values for weight or serum creatinine were given. In this case gentamicin toxicity was probably entirely predictable and therefore preventable.

The British National Formulary recommends that the dose interval for gentamicin should be increased to 12 hours when the creatinine clearance is 30-70 ml/min. Creatinine clearance falls with age, and a number of equations have been developed to predict this from age, sex, weight, and serum creatinine, assuming steady state fluid balance.1 All of these variables are available to the clinician before gentamicin is prescribed. According to the equation derived by Hull and others,1 an apparently normal serum creatinine of 90 µmol/l would give a creatinine clearance of greater than 70 ml/min only if the patient weighed more than 71 kg. Even at the upper end of the normal reference range a serum creatinine of 120 µmol/l would give a creatinine clearance of greater than 70 ml/min only if the patient were over 95-7 kg. The patient was probably prescribed a three times daily regimen on the basis that his renal function was normal because the serum creatinine value was within the normal reference range. This is a false premise, and in the elderly there can be important renal impairment with a normal serum creatinine concentration.

There is no place for the automatic prescription of gentamicin three times daily in the elderly. An estimate of creatinine clearance should be made using one of the equations available and the dose interval adjusted accordingly.

Zaske and others have shown that the elderly have wide variations in volume of distribution and elimination rates for gentamicin.2 It is important, therefore, to measure serum gentamicin values at least two or three times a week as the concentrations cannot be accurately predicted in spite of initial adjustments in dose and dosage interval.

Had the above factors been taken into account the patient would probably have been spared the symptoms of vestibular toxicity.

Michael Barnham
Michael McEvoy
Departments of Microbiology and Haematology
Harrogate General Hospital,
North Yorkshire HG2 7ND

Gonadotrophin hormone releasing analogues open new doors in cancer treatment

Sir,—Dr Jonathan Waxman (31 October, p 1084) states that depot preparations of gonadotrophin hormone releasing analogues may be an acceptable alternative to orchidectomy in the treatment of prostatic cancer.

Since 1982, 135 patients at Broadgreen Hospital, Liverpool, have undergone subcapsular orchidectomy as part of their treatment for prostatic cancer.1 In 100 this was done under local anaesthesia, general anaesthesia being required only for combined procedures. The average age was 77 years (range 48 to 98 years). Three patients refused orchidectomy. We have estimated the cost of treating these 135 patients with the luteinising hormone releasing hormone analogue goserelin if it had been available in 1982 assuming survival to be the same. The total cost would have been £293 000 and the cost in the last financial year £93 000. When hormone manipulation is indicated there is a cogent argument for the continued use of subcapsular orchidectomy. The costs of the operation are relatively low in National Health Service practice since the procedure requires neither general anaesthesia nor major theatre time and the recovery period is short. Patient acceptability in elderly men is high and the use of the term 'mutilating' in this context can only be considered emotive.

Unless luteinising hormone releasing hormone analogues are subsequently shown by controlled clinical trials to offer significant therapeutic benefits over orchidectomy then we suggest that their role in prostatic cancer should be limited to the 2-3% of patients who refuse orchidectomy.

A J Arnold
A D Desmond
Broadgreen Hospital,
Liverpool L14 3LB

AUTHORS' REPLY.—We appreciate Dr Swain's helpful reminder that gentamicin toxicity can in general be predicted and avoided. Our own interest in this case was in its neurological features; it seemed to us worth while to point out that severe gentamicin toxicity can be present with no cochlear symptoms and little in the way of obvious vestibular symptoms. We agree entirely that this underlines the need for identifying risk factors and monitoring serum concentrations at appropriate intervals.

We know from the patient's case record at the hospital where he was initially treated that he was seriously ill from a life threatening infection and that this and bacterial sensitivities governed the use of gentamicin. We hope that our case report and Dr Swain's comments increase doctors' awareness of the need for careful monitoring of gentamicin, especially when its use is required in the elderly patient.

Roderick Duncan
Ian D Melville
Institute of Neurological Sciences,
Southern General Hospital,
Glasgow G51 4TF

Waiting list statistics

Sir,—The finding of Dr A Lee, Mr B Don, and Dr M J Goldacre (7 November, p 1197) that 28% of patients on the surgical waiting list at any one time are found eventually not to have been admitted for their surgery is mirrored by similar findings at the time when they are offered admission. The first year's statistics compiled by the orthopaedic bed manager at the Leicester General Hospital show that overall 25% of patients offered admission do not accept; an up: about half electively cancel their admission and half do not attend, having failed to give any warning. The numbers vary from surgeon to surgeon and week to week, and though the failure rate is higher during the holiday months and at Christmas, there is an appreciable failure all through the year.

Dr Lee and others do not know the fate of those who were not admitted in the Oxford region, and it is not possible to estimate the extent of 'leakage' in my experience. The Oxford region has a resident population of 1·5 million people and a resident population of 45 million people in the Oxford Region, so we have to consider both the Oxford and the Oxford Region statistics. The Oxford Region has a resident population of 1·5 million people and a resident population of 45 million people in the Oxford Region, so we have to consider both the Oxford and the Oxford Region statistics.

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A D Desmond
Broadgreen Hospital,
Liverpool L14 3LB


References


Hull University, Hull HU3 2WY

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