Pneumonitis in an elderly Bangladeshi man

The incidence of primary varicella zoster virus infection (VZV) in young adults and pregnant women has risen in recent years and is accompanied by a greater risk of serious complications. VZV disease in the elderly usually presents as shingles, as a result of secondary reactivation of latent infection, and can be treated successfully with early antiviral therapy. We report a case of fatal primary infection in an elderly man.

A 66 year old Bangladeshi man with fibrosing alveolitis and non-insulin dependent diabetes mellitus was admitted to hospital with increasing shortness of breath for one week. He was a smoker and had been well controlled on 25 mg of prednisolone daily for the previous two months.

On admission he was febrile (38.5°C), tachypnoeic (50 breaths/minute), and hypertensive (blood pressure, 85/60 mm Hg), with severe mucosal candidiasis. An extensive maculopapular rash, present for three days, was noted and thought to be consistent with amoxicillin treatment, which had been started before admission. Coarse crepitations were heard throughout the chest, oxygen saturation was 80% on air, and blood gases showed type I respiratory failure (pH 7.3; partial CO2 pressure, 4.8 kPa; partial O2 pressure, 8.39 kPa; HCO3−, 19.4 mmol/litre). Haemoglobin was 180 g/litre, white blood cell count was 8.5 × 109/litre, serum creatinine was 180 mmol/litre, and blood glucose was 22.2 mmol/litre.

The chest x ray showed right mid zone consolidation (fig 1). Bronchopneumonia was diagnosed; intravenous ceftriaxone, clarithromycin, and fluconazole were commenced. After consultation with the virologists the patient also received aciclovir (10 mg/kg) intravenously.

He required intubation, ventilation, and inotropic support within 14 hours of admission. He remained persistently hypoxic despite 100% oxygen, positive end expiratory pressure, and ventilatory support. He remained persistently hypoxic despite 100% oxygen, positive end expiratory pressure, and ventilatory support. He remained persistently hypoxic despite 100% oxygen, positive end expiratory pressure, and ventilatory support.

..continued

Anaplastic large cell lymphoma: what's in a name?

We have read with interest the editorial by De Wolf-Peeters and Achten, which recently appeared in your journal. As the authors state in this editorial, the so-called anaplastic large cell lymphoma (ALCL) is probably more heterogeneous than currently recognised in the REAL/WHO classification.

Recent studies have shown the existence of a true clinicopathological entity among the CD30 positive anaplastic lymphomas; namely, an ALCL subtype characterised by a specific chromosomal aberration, involving the anaplastic lymphoma kinase (ALK) gene on 2p23, and by excellent prognosis.1 ALK expression is found in 30–60% (depending on the age of the population studied) of systemic ALCLs with (primary) nodal involvement. It is not found in primary cutaneous and other extranodal ALCLs.2,3

The remaining controversy concerns ALK negative systemic nodal ALCL, which may be morphologically indistinguishable from primary cutaneous ALCL, but which runs an aggressive clinical course compared with primary cutaneous ALCL. Furthermore, the distinction between ALK negative systemic ALCL and peripheral T cell lymphomas that otherwise specified (T-NOS) is drawn into question because the formerly acknowledged difference in prognosis between ALCL and peripheral T-NOS might result from the favourable prognosis of ALK positive ALCL. Thus, as the authors correctly state, criteria for proper distinction between (ALK negative) systemic ALCL and extranodal ALCL, as well as peripheral T-NOS, are needed to determine therapeutic strategies.

Recently, we identified a biological prognostic marker for ALK negative systemic ALCL.4 We found that a high percentage (>15%) of activated cytotoxic T lymphocytes (CTLs), present in the reactive infiltrate, is related to poor overall and progression free survival. This biological prognostic marker remained independent of, and seemed more sensitive than, established clinical parameters (such as the international prognostic index) in determining clinical outcome. The same relation between high numbers of activated CTLs and poor prognosis was found by our group for Hodgkin’s disease.5 These studies suggest that the interaction between tumour cells and reactive immune competent lymphocytes might be more important to clinical outcome than the morphology and immunophenotype of the lymphoma cells. As such, studies intending to clarify the distinction between the above

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When should a coroner’s inquest be held? The Manchester guidelines for pathologists

I am grateful to Dr Roberts and colleagues for their important paper and argument to formulate guidelines on when to report a death to the coroner and decisions thereafter. The guidelines have important implications for practitioners in primary and secondary care. One particular dilemma is when to discuss with the coroner what appears to be a natural death, but where the cause is unknown or where the practitioner, but has not consulted in the required preceding 14 days. This is compounded by the variation in attitudes of coroners and their officers to such discussions.

It is indeed a “grey area” where needed guidelines are required for doctors and coroners so that it can be determined more precisely when necropsies and inquests are required. To facilitate this process, death certificates and the second part of cremation forms could comprise specific additional questions. These should ascertain the extent to which the certifying practitioner, and the independent practitioner in the case of a cremation form (part 2; form C), are in agreement that the cause of death stated is beyond reasonable doubt, and so whether or not involvement of the coroner is required. In the case of cremation, where a body is permanently disposed of, the use of a second practitioner who discusses the case with the first practitioner incorporates a double check into the process. This check is supplemented by a third practitioner acting as the crematorium medical referee.

It is well known that most medical diagnoses are made through clinical history, examination, and investigations, so that in most situations unnecessary distressing necropsies and inquests could be avoided, although they may be required given the existing regulations.

Guidelines should be aimed at improving the specificity of the postmortem service as a research tool to be of benefit in three ways when the cause of death is unclear, namely: (1) educating the profession, (2) ascertaining where death is unnatural, and (3) most importantly, facilitating the grieving process of relatives. The latter could be facilitated by next of kin clinics, which have been discussed in this journal, and which target the needs of families attempting to come to terms with the complicating factors of a coronial investigation at a time of crisis, particularly in the situation of sudden death. Such changes should not only improve the quality of the service and its clinical effectiveness, but also be an aid to clinical governance in this area.

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Book reviews


Valproate has been the subject of over 5000 publications. It was discovered accidentally: it was used as a vehicle for some water insoluble compounds under investigation and found to have an independent anticonvulsant effect. The first clinical reports appeared in 1964. It can justifiably claim its place in the “milestones in drug therapy” series.

Valproate has been used in the treatment of epilepsy since the 1970s. In addition, it has been used in other conditions including bipolar affective disorder, chronic pain, and migraine. It has a wide range of side effects and has been associated with a number of adverse events. It is still one of the most prescribed anticonvulsant drugs in the world.

The first book is Valproate. Milestones in Drug Therapy. However, it is intended for specialist neurologists and is not suitable for a general audience. The second book is Valproate: Milestones in Drug Therapy. It is a comprehensive review of the scientific literature and provides an overview of the pharmacology, mechanisms of action, and clinical applications of valproate. The authors are predominantly from US centres, and the book is well written and extensively referenced. Anyone wishing information on valproate would be unlucky indeed not to find it within these covers.

If I have a criticism it would be that the editor has not been strict enough with his contributors. Thus—for example, the chapter on toxicity might be usefully aimed at doctors who are used to the chapter on side effects and the information on liver damage can be found in several places elsewhere in the book. Some contributions need pruning. The chapter on the use of valproate in children was twice as long as that in adults, which was fine except that much of the paediatric chapter was taken up with a description of different paediatric epilepsy syndromes; interesting and important information but not directly relevant. The length of the chapter on the use of sodium valproate in headache (20 pages) seems disproportionate in that there are only four double blind, placebo controlled trials in the use of this drug in migraine.

However, these minor criticisms do not detract from the value of this useful book, which should find its place in pharmacology departments and neurology libraries.

R. ROBINSON

Chemokines and Cancer. Rollins BJ, ed. ($125.00.) Humana Press, 1999. ISBN 0 89603562 X.

This multiauthor review has 16 chapters, broken down into sections on the physiology of chemokines, tumour infiltration by leucocytes, modulation of host response to cancer, chemokines and tumour growth and metastasis, associations with specific malignancies, and finally effects on stem cell proliferation. Authors are predominantly from US centres, with a scattering of Europeans.

I have to admit to finding this mostly hard going and, after starting off with good intentions and struggling through half of the book, it was laid aside as more easily digestible and clinically useful things usurped my attention. Half the problem is the multiplicity of chemokines, mostly with incomprehensible names, which makes it necessary for the non-specialist to keep referring back to the first chapter to have any idea about the remaining chapters. This is a fast moving field, so it is difficult to say what the value of this publication is except as a way mark for the long journey of discovery in cancer research.

There is little here for the working pathologist, but plenty for the chemokine and cancer immunology researcher. I doubt very much whether it will find a general market for interested non-specialists. Mainly one for the library.

G P SPIECKETT
Calendar of events

Full details of events to be included should be sent to Maggie Butler, Technical Editor, JCP, The Cedars, 36 Queen Street, Castle Hedingham, Essex CO9 3HA, UK; email: maggiebutler@pilotree.prestel.co.uk

Professional Standards of Pathologists in a Modern NHS Pathology Service
7 June 2001, Royal College of Pathologists, London, UK
Further details: Michelle Casey, Academic Activities Coordinator, 2 Carlton House Terrace, London SW1Y 5AF, UK. (Tel +44 020 7451 6700; fax +44 020 7451 6701; www.rcpath.org)

Recent Advances in Genetics
5 July 2001, Royal College of Pathologists, London, UK
Further details: Michelle Casey, Academic Activities Coordinator, 2 Carlton House Terrace, London SW1Y 5AF, UK. (Tel +44 020 7451 6700; fax +44 020 7451 6701; www.rcpath.org)

BSCC Annual Scientific Meeting
9–11 September 2001, Majestic Hotel, Harrogate, UK
Further details: BSCC Office, PO Box 352, Uxbridge UB10 9TX, UK. (Tel +44 01895 274020; fax +44 01895 274080; email lesley.couch@psilink.co.uk)

41st St Andrew’s Day Festival Symposium on Therapeutics
6–7 December 2001, Royal College of Physicians, Edinburgh, UK
Further details: Eileen Strawn, Symposium Coordinator. (Tel +44 0131 225 7324; fax +44 0131 220 4393; email 2.strawn@rcpe.ac.uk; website www.rcpe.ac.uk)

Current Concepts in Surgical Pathology
12–16 November 2001, The Four Seasons Hotel, Boston, Massachusetts, USA
Further details: Department of Continuing Education, Harvard Medical School, PO Box 825, Boston, MA 02117-0825. (Tel +1 617 432 1525; Fax +1 617 432 1562; email hms-cme@harvard.edu; web page http://www.med.harvard.edu/conted/)
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