Aetiology of chronic suppurative lung disease

EDITORS—Nikolazik and Warner report that eight of 41 children with suppurative lung disease had a demonstrable immunodeficiency.1 Four of these patients were shown to have abnormalities of neutrophil function on the basis of some form of in vitro functional assay. Three further patients with other causes for their lung disease were said to have similar defects.

None of these deficiencies are recognised within the comprehensive World Health Organisation classification of immunodeficiencies. This is probably because such functional abnormalities have proved difficult to characterise in a consistent manner. The possibility exists that the observations in these assays are due to immunological activation secondary to chronic infection.

I am surprised at the relative absence of immunoglobulin and antibody deficiencies in this series. These are the most common primary immunodeficiencies and are well characterised as leading to suppurative lung disease. A proportion of the 37% of children without identifiable predisposing factors might have such deficiencies if they had detailed investigation of their immunoglobulin and antibody production. The methods section does not outline the extent of the immunological evaluation or whether all patients received such an evaluation. The identification of such patients is important as replacement immunoglobulin treatment may be the only way of preventing progression of their pulmonary disease.

Finally the two patients with measles and adenosine infection and measles and disseminated Mycobacterium intracellulare infection are placed in the secondary immunodeficiency group. These patients may also have primary cell mediated immunodeficiency, and indeed, the reader is left with the impression of a severe and complex course of measles is often associated with a primary immunodeficiency.

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Professor Warner comments:

Dr Morgan makes a number of important points about our publication that certainly require answering. The presentation of a short report does rather restrict the number of references that can be given which might perhaps have provided some of the quality and quantity Dr Morgan is concerned that the deficiencies described are not recognised within the World Health Organisation classification of immunodeficiencies. This classification is very exclusive. The criteria are very restrictive and this is rather like placing blinkers on immunologists in suggesting that significant and reproducible abnormalities of immune responsiveness are not evidence of immunodeficiency. This is patently not the case. It is important to emphasise that none of the immune investigations were done when children had active infection. The C reactive protein was always normal at the time of investigation, otherwise the results were rejected. Furthermore, the vast majority had more than one investigation to confirm the abnormality. I would refer Dr Morgan to our publication which was reference 6 in the original paper.1 This describes the neutrophil defects in more detail. It also provides extra evidence which might eventually allow them to be recognised within the classification system.

Sadly most of the investigation in these children was done at a time before there was a common awareness of IgG subclass deficiencies. Therefore, this was a notable shortcoming, and I do accept that we should perhaps have identified some immunoglobulin and antibody deficiencies. However, the presence of IgG subclass deficiency remains a very difficult area where there are often explanations for bronchiectasis. In this respect, two of the patients with primary ciliary dyskinesia syndrome had neutrophil defects. Dr Morgan might be interested in knowing of two studies which have been done suggesting that there might be primary defects of neutrophil function in patients with this syndrome which may relate to abnormalities in cytoskeletal proteins.3 The patients with measles all had their immune functions studied long after recovery from measles and the responses were normal. We certainly had no evidence of any abnormality of cellular immune response at the time that they were investigated for suppurative lung disease. This, of course, does not totally exclude the possibility that there was a rather specific susceptibility to measles, though the severity of the measles’ illnesses in each case was not consistent with this. The association between measles and secondary adenosine infection is well described.


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Professor Warner comments:

Sedation for investigations: prolonged effect of chloral and trimeprazine

EDITORS—Sedation is increasingly required for procedures such as computed tomography which are not painful but require the child to remain still during the investigation. Young children in particular may need to be heavily sedated to achieve satisfactory results. We have performed a survey of the use of sedation for investigations in our paediatric department. This included telephone follow up of children after discharge.

Data were obtained for 34 children, median age 2.8 years. Procedures included 17 computed tomograms and 12 auditory evoked response studies. The drugs most commonly prescribed were chloral (31 patients, median dose 50 mg/kg) and trimeprazine (32 patients, median dose 2.9 mg/kg). Seventeen procedures were recorded as easy to perform, 12 were performed with difficulty, and five failed. Ward staff commented on problems relating to sedative medication in 18 children. Two were too drowsy for discharge from the day case unit and required admission overnight. We also audited the use of chloral sedation and the use of monitoring. In general these fell below standards that would be considered safe in children having general anaesthesia, despite the use of heavy sedation.

Anesthetists should have a central role with paediatricians and radiologists in the development of sedation protocols for children. We have found that the use of chloral and trimeprazine for sedation of children is associated with prolonged drowsiness. Standardisation of monitoring and post-anaesthetic care in children requiring heavy sedation should be similar for those having general anaesthesia.


Developmental setback in severe visual impairment

EDITORS—The data presented by Dr Cass and her colleagues are of importance and significance for paediatricians and other
Clinical details of six children studied

<table>
<thead>
<tr>
<th>Year of birth</th>
<th>Ophthalmological diagnosis</th>
<th>Period of normal development (years)</th>
<th>Perceived environmental stress factors</th>
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<td>Anophthalmia</td>
<td>3-3-5</td>
<td>Hospitalisation</td>
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<tr>
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<td>Hypoplastic retinal membrane</td>
<td>2-3-25</td>
<td>Separation from parents</td>
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<td>2-2-5</td>
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<tr>
<td>M 1979</td>
<td>Optic nerve hypoplasia*</td>
<td>2-5-3</td>
<td>Nursery placement, birth of brother</td>
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<tr>
<td>M 1982</td>
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<td>2-5</td>
<td>Nursery placement</td>
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<tr>
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<td>Ocular haemangiomata</td>
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*Light perception present. All six are now severely retarded with overt autistic behaviour.

professionals who are concerned with assessing and attempting to predict the developmental outlook for children with severe visual handicap. The authors make clear that it is those children with the most severe degrees of visual impairment who run the greatest risk of sustaining developmental setbacks and suggest that component causative factors include primary maldevelopment of the central nervous system, the degree of visual impairment, the developmental stages that affected children have reached and the developmental and emotional environments to which they are subjected.

The developmental profiles of some of the children they describe are reminiscent of children who show the features of disintegrative psychosis in childhood, in which it may be that inherently vulnerable children develop a severe acquired autistic syndrome often in association with evidence of severe environmental stress. A similar picture can be seen in children with severe visual impairment and over recent years I have followed up the progress of six children of whom had severe visual impairment and who have shown evidence of severe and permanent developmental setbacks. Their clinical details are summarised in the table.

Clearly, the experiences of Cass and her colleagues are likely to represent not uncommon phenomena. I would endorse their recommendations that careful attention be given to the developmental and environmental vulnerability of this group of handicapped children, particularly as the developmental setbacks that they describe are likely to cause major and long term disability.

Ambulatory paediatrics

EDITOR.—The paper by Doug Heller gave an excellent overview of the philosophy of ambulatory paediatrics. I agree that it is a change in attitude, making the service more child and family oriented, rather than a new specialty.

The definition of ambulatory paediatrics is broad, and many people find the concept a little abstract. I would like to give concrete examples of two successful developments within our ambulatory service.

A daily urgent consultant led clinic has been introduced. Many similar clinics suffer from being overcrowded by inappropriate referrals. We have avoided this by introducing a ‘hotline’, which is a direct line manned by a consultant and runs for one hour before the urgent clinic. General practitioners can phone for advice and to discuss patients. Those thought to need an urgent appointment are seen that day. Preliminary data suggest that less than half the calls to the hotline result in an urgent appointment, and many problems can be dealt with by advice only.

The second development concerns our children’s home care nurse who rotates through the accident and emergency department. The aim was to improve the quality of care for children in the accident and emergency department and to reduce short stay admissions. The initiative was initially supported by and is currently being evaluated by the King’s Fund. Preliminary data suggest this has been very successful, and we are committed to continuing the service.

Role of ultrasound in congenital hip dysplasia

EDITOR.—We were pleased that the publication of the annotation on this subject in the April issue gave prominence to the potential of this approach to the management of babies with clinically suspected hip instability. We are convinced that paediatrics is an exciting and challenging area to be involved in, with much scope for development. Even small changes in attitude and in the way the service is delivered can make a huge difference to the quality of care provided.

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