NSF for children and young people in Wales

The National Service Framework (NSF) for children, young people, and maternity services in England has recently been published by the Department of Health. On 15 October the equivalent Welsh consultation document was launched by the Minister for Health and Social Services at the Welsh Assembly Government, Mrs Jane Hutt; the consultation document of the Welsh NSF differs from the English NSF in a number of ways.

First, the Welsh document has set standards not just for health and social care but also for the other agencies which have a strong influence on the health and wellbeing of children, such as education, housing, leisure, and transport. The document has been endorsed in a foreword signed by the First Minister, Rhodri Morgan, and all members of the Cabinet of the Welsh Assembly Government.

A second difference from the English NSF is that the Welsh document is written with 21 standards and 203 “key actions”, each of which is specific, written in the present tense, and can be measured. The third difference is that a web based self-assessment audit tool is being developed for publication at the same time as the final standards in the summer of 2005, which we hope will enable local measurement of progress in achieving the key actions. In addition, the standards and key actions will be subject to inspection processes by Health Inspectorate Wales (HIW) (working in conjunction with the Health and Social Commission), and we anticipate that there will be joint inspections carried out in Wales between HIW, Social Services Inspectorate Wales, Care Standards Inspectorate Wales, and the Inspectorate for Education and Training in Wales (ESTYN).

It is clear that the profile of the children’s and young people health and wellbeing is higher than it ever has been on the political and planning agenda, in both England and Wales. We can learn from each other and we, in Wales, would be grateful to receive comments from all parts of the UK on our Welsh NSF, as part of the consultation process, before the final document is published in summer 2005. Please access the website www.wales.nhs.uk/nsf and let us have your comments.

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Competing interests: none declared

Risk of life threatening apnoea after immunisation

It is common practice in the UK to immunise babies with DTP (diphtheria, pertussis, tetanus), Hib (haemophilus influenza type b), meningitis C, and polio at 8 weeks following delivery, regardless of corrected gestational age. However, such recommendations may be inappropriate for premature babies who may be at increased risk of apnoea and bradycardia.

Twenty seven week gestation twins underwent their first immunisations at 62 days of age (corrected age 16.5 weeks). The following morning (15–16 hours later) both twins went a pale, dusky colour and had a respiratory arrest requiring bag and mask ventilation with severe bradycardia. On examination both infants were tachypnoeic and pyrexial. Intravenous antibiotics were started but stopped 48 hours later when full septic screens, viral cultures, and a nasopharyngeal aspirate returned negative. Cardiac and oxygen saturation monitoring showed several further episodes of desaturations requiring facial oxygen and gentle stimulation. A full recovery to normal self-ventilation in air followed over four hours. Both babies remained pink, active, and with no respiratory distress despite elevated C reactive protein (twin 1: 27 mg/l; twin 2: 38 mg/l; normal <10 mg/l) and raised platelets in twin 1 (420 x 10^9/l; normal 190-400). The reactions were reported to the UK Adverse Drug Reactions reporting scheme.

Four weeks later the second immunisation sets using acellular pertussis were administered while cardiorespiratory and oxygen monitoring was performed. Neither twin had any reaction to the second course of immunisations. Current evidence points to an increase in episodes of apnoea and bradycardia in pre-term infants receiving their eight week immunisation, and the unit has decided to review its policy on the monitoring of such infants.

The episodes of apnoea and bradycardia in the twins following their immunisations were highly suggestive of a delayed type hypersensitivity reaction to a component of one of the vaccines. Studies have implicated the whole cell pertussis component of DTwP with significantly more reactions and raised C reactive protein after immunisation with DTwP than after separate diphtheria, Hib, and tetanus toxoid vaccines alone.

Current opinion for immunisation of pre-term infants suggests cardiorespiratory monitoring for up to 48 hours post-immunisation rather than postponement of immunisations; however, recommendations for future immunisations in infants who have had an episode of apnoea or bradycardia are unclear. Many suggest immunisation with acellular pertussis, as inpatients with 48 hour monitoring. However, the evidence from one Australian study implies that the risk of future reaction is very low.

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doi: 10.1136/adc.2004.054544
Competing interests: none declared

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3 Anderson CP, Fineberg HV. Adverse events following pertussis and rubella vaccines. JAMA 1992;267:392–6.
Addressing child welfare concerns: a new approach

In a small proportion of childhood hospital attendances there are obvious child protection issues. In a much greater number there is concern about the child’s welfare. Only if concerns are recognized, documented, and addressed at an earlier stage, can we hope to improve “safeguarding” children.

In Peterborough District Hospital a “Concern Sheet” has been in use since 1998 to address child protection concerns throughout the Trust. Use of the Concern Sheet has been audited twice. Despite this, many failures of documentation, reporting, and follow up were identified. To address these problems, a joint hospital/community “Children’s Liaison and Discharge Coordinator” was appointed in October 2002. She is a registered children’s nurse, with child protection experience.

Our aim was to see if these two measures improved identification, documentation, and follow up of child protection concerns. We retrospectively analysed the Concern Sheet data collected for 2003 and noted a striking increase in child protection awareness in every hospital department where children are seen (table 1). The Coordinator has been very active in raising the profile of child welfare concerns, not just over abuse, with all staff.

Table 2 shows the causes of concern for different age groups. It is interesting that 25% of reported concerns were about parents and their ability to care for their children. There is a potential risk to children cared for by adults with mental health problems, those who abuse drugs/alcohol, or when there are concerns regarding domestic violence. Hall has stressed that healthcare professionals must take the opportunity to prevent child abuse/neglect when faced with such situations.

We believed that we were addressing concerns which were less serious at an earlier stage, but it is noteworthy only 47 (9.6%) required no further action. Nearly half (230, 46.9%) of the concerns were serious enough to warrant a discussion with Social Services. The Coordinator liaised with health visitors in 229 (46.7%), school nurses in 21 (4.3%), Child and Adolescent Mental Health Services in 29 (5.9%), and police in 40 (8.2%) of the cases where Concern Sheets were completed.

Forty two children (18.2%) proceeded to an Initial Child Protection Conference, 14 (6%) had an early Review Conference, and 2 (0.8%) had an early Transfer-in Conference as a result of the concern reports. Of those subjected to an Initial Conference, 36 (86%) were registered.

From past enquiries into child deaths, the common reasons which have led to a failure to intervene early enough are poor training, documentation, information sharing, and follow up of concerns. Lord Laming’s report has emphasised the importance of better training and introducing systems which allow quality monitoring. We have attempted to address these issues and conclude that “safeguarding children” may be improved by:

- Having a person other than the named and designated professionals in the role of a Coordinator
- Having a uniform way of recording child welfare concerns throughout a Trust.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Areas of the Trust and numbers of Concern Sheets compared for the years 2002 and 2003</th>
</tr>
</thead>
<tbody>
<tr>
<td>Area</td>
<td>2002 (n = 153)</td>
</tr>
<tr>
<td>Paediatric assessment unit</td>
<td>68 (44.4%)</td>
</tr>
<tr>
<td>Paediatric ward</td>
<td>43 (28.1%)</td>
</tr>
<tr>
<td>A&amp;E</td>
<td>32 (20.9%)</td>
</tr>
<tr>
<td>Special care baby unit</td>
<td>3 (1.9%)</td>
</tr>
<tr>
<td>Children’s outpatients</td>
<td>3 (1.9%)</td>
</tr>
<tr>
<td>Community child health</td>
<td>0</td>
</tr>
<tr>
<td>Maternity unit</td>
<td>0</td>
</tr>
<tr>
<td>Others*</td>
<td>4 (2.6%)</td>
</tr>
</tbody>
</table>

*As awareness increased, concerns were also received from surgical wards, therapists, and paramedics in 2003.

<table>
<thead>
<tr>
<th>Table 2</th>
<th>Concerns categorised according to age group</th>
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<tbody>
<tr>
<td>Category</td>
<td>0–4 years (n = 260)</td>
</tr>
<tr>
<td>Neglect</td>
<td>102 (39.2%)</td>
</tr>
<tr>
<td>Physical abuse</td>
<td>94 (36.1%)</td>
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<tr>
<td>Sexual abuse</td>
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</tr>
<tr>
<td>Emotional abuse</td>
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<tr>
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</tr>
<tr>
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<td>0</td>
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<tr>
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Each column adds up to 100%.

References

Maternal vitamin D deficiency, refractory neonatal hypocalcaemia, and nutritional rickets

We read with interest the articles by Allgrove’ and Ladhani and colleagues’ which highlighted the re-emergence of vitamin D deficiency, neonatal hypocalcaemia, and nutritional rickets. We would like to present our experience from a single centre of maternal vitamin D deficiency, neonatal hypocalcaemia, and nutritional rickets. We believe that we were addressing concerns which were less serious at an earlier stage, but it is noteworthy only 47 (9.6%) required no further action. Nearly half (230, 46.9%) of the concerns were serious enough to warrant a discussion with Social Services. The Coordinator liaised with health visitors in 229 (46.7%), school nurses in 21 (4.3%), Child and Adolescent Mental Health Services in 29 (5.9%), and police in 40 (8.2%) of the cases where Concern Sheets were completed.

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References

Maternal vitamin D deficiency, refractory neonatal hypocalcaemia, and nutritional rickets

We read with interest the articles by Allgrove’ and Ladhani and colleagues’ which highlighted the re-emergence of vitamin D deficiency, neonatal hypocalcaemia, and nutritional rickets. Leicester City has an estimated proportion of 28% South Asians (Census 2001) and an increasing number of other ethnic groups, including an estimated recently arrived 10 000 people of Somalian origin. Studies in our centre have confirmed that significant numbers of south Asian mothers have vitamin D deficiency at the end of pregnancy, and substantial numbers of children have infantile and adolescent rickets, some of whom have extremely severe bony deformities. In addition there have been increasing numbers of late (5–10 days of age) and late-late (2–12 weeks of age) neonatal hypocalcaemia, presenting predominantly with seizures, which, despite intensive calcium and vitamin D treatment have been difficult to correct biochemically. All the mothers had vitamin D deficiency and were supplemented with oral vitamin D and calcium supplements. None of the mothers, despite being within high risk ethnic groups, had vitamin D supplementation in pregnancy despite the recommendation by COMA (Committee on Medical Aspects of Food Policy in UK) that all at-risk pregnant and lactating mothers should receive 10 μg (400 IU) of vitamin D daily. Furthermore, a local audit involving clinicians in antenatal care including general practitioners, midwives, and obstetricians showed that, while health professionals were aware of this issue, there was no clear policy followed.4

At birth, the newborn’s vitamin D status is directly related to maternal vitamin D status and maternal-fetal transfer of vitamin D and its metabolites in pregnancy. Babies whose mothers have a marked vitamin D deficiency will have a compromised vitamin D status,4 and this has important long term implications for the health of the offspring.4

As noted by Allgrove there were national and local “stop rites campaigns” in the 1970s, and in Leicester this appeared to reduce but not remove the spectre of nutritional rickets. In view of our recent experiences confirming an increasing frequency and severity of neonatal vitamin D deficiencies, vitamin D supplementation in addition to calcium is advised.3

www.archdischild.com
deficiency we would strongly agree with Allgrove and Ladhani and colleagues in emphasising the importance of vitamin D supplementation. It is certainly a serious indictment of our community preventative services not to have protected “high risk” mothers and their offspring. We would propose an urgent review and implementation of the national recommendations on vitamin D supplementation in “high risk” pregnant women and infants to prevent associated infantile co-morbidity.

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References

Apnoeas in bronchiolitis: is there a role for caffeine?

Apnoeas in bronchiolitis is a common respiratory illness in infants in winter months. Recurrent apnoeas in high risk infants with severe bronchiolitis increases the need for respiratory support (nasal continuous positive airway pressure and ventilation) and transfer to the paediatric intensive care unit (PICU). During the winter of 2003–04 we had three babies presenting with apnoeas secondary to bronchiolitis. All three babies were ex-premature infants under 3 months of age. All had deterioration in their respiratory status potentially needing further care in PICU. On advice of two PICU consultants these babies were treated with a loading dose of caffeine. All three showed immediate improvement in their respiratory status and avoided being transferred out. Caffeine is a respiratory stimulant widely used in the treatment of apnoea of prematurity. Following our experience we performed a questionnaire survey of the use of caffeine for apnoeas in bronchiolitis across 20 intensive care units in the UK. We made a thorough literature search to look at the evidence.

Of the 20 questionnaires sent out, only 10 replies were received. Opinion was divided between PICU consultants, with four stating that they would advise a trial of caffeine. This made a total of six, including the two who advised us previously. The evidence from literature is anecdotal.

We conclude that there is little evidence in literature to support the use of caffeine in bronchiolitis, and there is divided opinion in PICUs across the UK. We feel that caffeine is a relatively simple treatment option in a district general hospital for apnoeas in bronchiolitis and recommend a randomised controlled trial. We would welcome comments on similar experiences from readers.

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doi: 10.1136/adc.2004.064824

Competing interests: none declared

BOOK REVIEWS

Child public health


R Tomlinson, S Lenton


Management of epilepsy in children can be complex and challenging, and a good clinician knows when to draw on multidisciplinary professional expertise, while staying up to date with clinical and non-clinical areas outside his or her immediate expertise. No one understood this more than the late Sheila Stella. The next three chapters give an excellent summary of the theories, key concepts, and techniques used in child public health.
train in the 1980s. She was a hands on clinician who did not shy away from basic science. The first edition of her book, *Epilepsy in children*, published in 1996, not surprisingly encompassed the various disciplines involved in the understanding and management of epilepsy, and became a popular reference text. When I picked up the second edition, now edited by Sheila Wallace and Kevin Farrell, I knew the book would be good reading but would it be better and worth shelling out another £120.00? The short answer is yes. As in the first edition this book covers all aspects of epilepsy in children with additional contributing authors reflecting the internation scope. There is also a distinct change in style to include the modern trend of boxes with key points at the end of each chapter, and tables allowing brevity of text. At times there is a price to pay for the brevity. For example I was disappointed that the initial chapter did not include the old classification of seizures and epilepsy syndromes, when the new classification is only a proposal and authors in the text continue to use the old terminology. Similarly the chapter on chromosomal syndromes only touched on disorders, even when epilepsy is the major presentation, such as in ring chromosome 20. There are however many chapters of substance covering pathology, pathophysiology, neuropsychology, and neuroimaging, in addition to the various age dependent epilepsy syndromes and lesional epilepsies including following brain injury. The chapter on psychiatric and cognitive aspects of epilepsy, areas often neglected or left to other professionals, are a must reading for paediatricians developing an interest in epilepsy and the more experienced clinician. Treatment of epilepsy including surgery and management of status epilepticus is well covered, though for the UK reader it is unfortunate that while the ketogenic diet is included, there is no mention of the modified ketogenic diet offered in our centres.

In the first edition Sheila Wallace tackled the chapter on neurophysiology herself, but rightly invited neuropathologists to contribute for this edition. These authors provide an excellent overview of the normal and abnormal EEG patterns for the clinician with super clarity of EEG samples included with the text. The chapter would have been enhanced by clearer guidelines on the use of the EEG, especially the value, if any, of repeat interictal EEG recordings in the management of children with epilepsy, covering issues related to drug withdrawal, cognitive deterioration, etc. Though the EEG is a valuable and relatively inexpensive tool, its limitation is not always appreciated.

With the proposed establishment of epilepsy networks in the UK and the expectation for a named paediatrician in each district general hospital with an interest in epilepsy, this book is well placed to be a valuable addition to the various pediatric libraries and also an informative reference source for the paediatric neurologist.

Z Zaiwalla

Recent advances in paediatrics 21


The latest in the Recent advances in paediatrics series intends, as the preface states, to provide a review of important topics and help keep doctors abreast of developments in the subject. It is aimed at practising clinicians, trainee paediatricians, and those preparing for specialty examinations. It contains 14 chapters covering a variety of general paediatric, neonatal, and community paediatric topics, as well as a literature review listing key articles and selected reviews published in 2002. The chapters themselves are generally broken down into specific areas for debate, and round off with a listing of key points for clinical practice and a literature review.

The subject matter chosen is varied and diverse, including summaries of recent developments and current practise in areas such as Kawasaki disease, asthma, diabetes mellitus, idiopathic thrombocytopenic purpura, inhaled nitric oxide in the newborn, the use of cannabis in teenagers, and childhood depression. The chapters are well written by respected authors in the appropriate fields. They break down well into bite sized chunks of easily digestible information, and contain a good sprinkling of diagrams and tables, with the occasional radiograph and clinical photograph. There are excellent treatises on very common and relevant areas in which the literature is traditionally rather neglectful. The chapters on head lice, cannabis use, weaning from assisted ventilation, and safe sedation provide invaluable advice and experience in dealing with everyday clinical situations on which little is generally written. The “Key points for clinical practice” boxes provide wonderful, concise summaries of the preceding chapters, although one feels that it may have been more effective to keep each box restricted to one page of the book instead of frequently spilling over into two. The diagrams and photographs are relevant, but occasionally a little fuzzy and sadly lacking in colour. The radiographs and photographs are clear, but seem on the whole to add little to the subjects. The literature review is very well set out and would be useful for further reading.

The book is aimed at clinicians and trainees and does indeed cover topics in general and community paediatrics and neonatology. It will probably be more relevant to non-specialist practitioners and more junior trainees aiming to update their knowledge, but is unlikely to be a substantial enough review for those in more specialist areas such as neonatology. It may not add much to the cause of passionate specialisations other than to direct further reading. Where this book will absolutely shine, however, is as a teaching aid. Whether it is used as a reference for common topics or as summary of the recent literature, it will provoke discussion and debate and is likely to engage the reader in the pursuit of further knowledge. It could serve well as the basis for a series of journal clubs or a starting point for the development of departmental protocols. Even better, it could be used in directed small group teaching with medical students, senior house officers, GP trainees, or core paediatric specialist registrars in order to summarise current opinion, promote the exchange of ideas and experience, and guide further reading and study. Not every clinician, but certainly every paediatric department could make excellent use of this book.

L R Wisby

The clinical management of craniosynostosis


In the introduction of this text the editors state that they have individualised their desire to produce a volume that does not constitute a specialised text for experts, but rather a source of information for associated professionals who perform an essential role in the management of these complex problems, at a significant distance from the “home institution”. In this regard the book has been successful in its aim, and it is certainly refreshing to see a book of craniofacial surgery not concentrating on surgery for craniosynostosis.

In fact the contents of the book highlights the fact that surgical intervention, although a key factor in the treatment of these complex cases, is only a relatively small/short phase of a coordinated multidisciplinary care pathway in these frequently complex cases. The authors have acknowledged the fact that this text constitutes a single unit, a philosophy, and as such is a distillation of the various roles to which a major busy unit with a substantial case load.

These are presented in clear sections supported by good literature reviews. As with any texts from multiple authors overlapping in terms of clinical expertise, the book does reflect a degree of repetition, particularly in the earlier chapters.

The overall quality of sections is good, with some excellent chapters, particularly on ophthalmology, airway management, and issues addressing raised intracranial pressure. Certain sections which are highly complex may lose the non-specialist readers. These are mainly related to the genetics and particularly the aetiopathogenesis of the conditions. This is a complex subject which realistically is of such a nature that it cannot effectively be simplified without becoming meaningless.

As indicated by the authors, the surgical options presented are not meant as a surgical atlas but are aimed at giving an overview of the options possible. They have acknowledged the fact that the surgical techniques presented are a number of many options varied by individual preferences. This specific chapter is one which could possibly have benefited from photographic illustrations in pre- and post-operative cases, rather than relying on simple line diagrams as the only way of presentation.

In conclusion, this text represents a worthwhile contribution to the craniofacial literature. It is generally a readable and accessible source of information, achieving the aims outlined by the editors and all contributors should be congratulated on a book highlighting the fact that a coordinated multidisciplinary approach is essential in the treatment of all patients with craniosynostosis.

S A Wall