Towards evidence based medicine for paediatricians

Edited by Bob Phillips

Putting evidence into practice: part 2

How do you remember what you read in a paper? How do you call to mind that reference about petechiae and vomiting that means you don’t have to stick this perfectly well 8 year old needle-phobic? Capturing the outputs of your critical appraisals so you can retrieve them is easier if you formally record them in some way. These records have been described by lots of different people in lots of different ways. They are called a number of things, mostly with punishingly painful acronyms attached. Hence you get “critically appraised topics” (CATs—probably the most generic phrase), “best evidence topics” (BESTs), or “patient orientated evidence that really matters” (POEMS) depending on where you live or work. The topics you read here in Archimedes are printed as BESTs, with fuller individual CATs on the ADC website. A template you can fill in with your own questions is also available on the website, along with Instructions to Authors.

The underlying principles of all these different records are fairly similar. They have a headline which describes either the question under consideration or a compact form of the answer. They have the question that triggered the CAT to be created. They detail the study methodology and comment on strengths and weaknesses, and relay the relevant results. They may have further comments, but nearly all will have a summary of the clinical bottom lines.

Once you’ve created your summaries, you still need to keep them accessible. Placing them all (with copies of Archimedes) in a big red folder might seem a bit cheesy, but it’s been shown to be an effective way of getting clinical questions answered quickly.1 It also doesn’t rely on your hospital computer working, and can be safely accessed while drinking coffee. It has the benefit of having instant feedback too—with the aid of a pencil or pen, comments can be added to the margins without difficulty. The more technologically minded might want to put your own on a website. Many widely used word processing programmes will allow you to save your version of an evidence summary as a web (html) page. (From there, getting them onto a website is then a case of finding someone with a bit of IT knowledge.) Developing the technological idea further, a PDA (handheld computer) version could be made by those with adequate skills.

Of course, the next logical step when you’ve worked with your colleagues to produce a high quality summary of the evidence related to a clinically important question is to submit it to Archimedes. You too could be the next proud provider of information which makes clinical paediatrics more evidence based.

Reference

Short stature and the probability of coeliac disease, in the absence of gastrointestinal symptoms

Report by
J C W van Rijn, F K Grote, W Oostdijk, JM Wit,
Department of Paediatrics, Leiden University Medical Center, Netherlands; f.k.grote@lumc.nl
doi: 10.1136/adc.2004.057851

A 8 year old girl is referred to the paediatrician because of her stunted growth. Her height SDS is -2.3 (=1st centile); her weight for height ratio is +0.9 SDS (=80th centile). At the age of 3 years, her height SDS was +0.3 (=60th centile). Her weight for height ratio had not changed considerably. At the age of 4–5 years, she had several episodes of constipation and anaemia. There were no other complaints. A diagnostic work-up was performed: IgA anti-endomysium (IgA EmA) antibodies were positive. A small intestinal biopsy (gold standard) showed total villous atrophy, consistent with coeliac disease. During follow up, she fulfilled the ESPGAN criteria for coeliac disease (finding of characteristic small bowel mucosa abnormalities in a small bowel biopsy, and clinical remission when placed on a gluten-free diet).

Structured clinical question
What is the prevalence of coeliac disease [outcome] in children with short stature and no gastrointestinal symptoms [patients]?

Search strategy and outcome
Pubmed—(body height AND (short OR little OR small OR abnormal) OR body stature OR dwarfish) OR failure to thrive AND coeliac disease (limited by: Ages: All child 0–18 year, Language: English);120 references of which 11 relevant and of sufficient quality (see table 1).

Embase—same search strategy; no additional relevant references.

Cochrane database—same search strategy; none relevant.

Additional information on each of the topics is available on the ADC website (www.archdischild.com/supplemental)

Clinical bottom line
- In 2–8% of children with short stature and no gastrointestinal symptoms, coeliac disease may be the underlying cause.
- Excluding other causes for short stature increases the risk of having coeliac disease by 19–59%.
- Children with short stature should be evaluated for coeliac disease.

Acknowledgements
The financial support of Pfizer is greatly acknowledged.

REFERENCES

www.archdischild.com
There is an association between idiopathic short stature and coeliac disease.

### Table 1: Relation of short stature and coeliac disease

<table>
<thead>
<tr>
<th>Citation, country</th>
<th>Study group</th>
<th>Study type (level of evidence)</th>
<th>Outcome</th>
<th>Key results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Knudtzon et al (1991), Norway</td>
<td>168 children (50 girls; 93 boys; age 0.5–17.2 years) with short stature without significant abdominal symptoms</td>
<td>Prospective cohort study (level 2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>2.9% (5/168) of the children with short stature were diagnosed as having coeliac disease</td>
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<td>Stenhammer et al (1986), Sweden</td>
<td>87 children (32 girls; 55 boys; age 1.0–16.5 years) with short stature (height more than 2SD below the mean for age and sex) and no gastrointestinal symptoms, signs of systemic disease or malabsorption</td>
<td>Prospective cohort study (level 1b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>5% (4/87) children with short stature were diagnosed as having coeliac disease</td>
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<td>Cacciari et al (1983), Italy</td>
<td>108 patients (30 girls; 78 boys; age 2.8–16.7 years) with short stature (height below third centile) and no gastrointestinal symptoms</td>
<td>Prospective cohort study (level 1b/2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>8.3% (9/108) patients with short stature were diagnosed as having coeliac disease</td>
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<td>Cacciari et al (1983), Italy</td>
<td>60 children (21 girls; 39 boys) with short stature (height below third centile) and no gastrointestinal symptoms</td>
<td>Prospective cohort study (level 1b/2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>8.3% (5/60) patients with short stature were diagnosed as having coeliac disease</td>
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<td>Rossi et al (1993), USA</td>
<td>117 children (age: 2–17 years) with height more than 2SD below the mean for age. Of these children, 57 were diagnosed with GH deficiency. All children were clinically and chemically euthyroid</td>
<td>Prospective cohort study (level 2b)</td>
<td>Proportion of coeliac disease in a group of children with short stature</td>
<td>1.7% (2/117) of children with short stature had biopsy proven coeliac disease</td>
</tr>
<tr>
<td>Bonamico et al (1992), Italy</td>
<td>49 children (27 girls, 22 boys; mean age 112 months (SD: 39)) with short stature (height below the third centile) and no gastrointestinal symptoms. None of the 49 patients showed somatic, cardiac, renal or chromosomal disorders</td>
<td>Prospective cohort study (level 1b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>59.1% (29/49) children with short stature were diagnosed as having coeliac disease</td>
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<td>Groll et al (1980), UK</td>
<td>34 children (16 girls, 18 boys; age 2.5–17.0 years) with short stature (more than 2SD below the mean for age) and no gastrointestinal symptoms. There were no dysmorphic features, and endocrine investigations were normal</td>
<td>Prospective cohort study (level 1b/2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>21% (8/34) children with short stature were diagnosed as having coeliac disease</td>
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<td>Rosenbach et al (1986), Israel</td>
<td>23 children (12 girls, 11 boys; age 6–16 years) below third centile for age and a bone age delay of at least 25%. Extensive preliminary work up (including hypothalamic, pituitary, adrenal, and gonadal functions, sweat test, stool examination for ova and parasites) was found to be negative</td>
<td>Prospective cohort study (level 2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>48.7% (11/23) of the patients with short stature were diagnosed as having coeliac disease</td>
</tr>
<tr>
<td>de Lecea et al (1996), Spain</td>
<td>118 children (49 girls, 69 boys; age 11 months to 14 years), with height less than third centile for age. Preliminary work up (absorption, hormonal and genetic studies, sweat test, x ray for bone age, serum IgA, AGA) was performed</td>
<td>Prospective cohort study (level 2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>18.6% (22/118) of children with short stature had biopsy proven coeliac disease</td>
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</tbody>
</table>
Is an intravenous fluid bolus of albumin or normal saline beneficial in the treatment of metabolic acidosis in a normovolaemic newborn?

Report by Hassib Narchi, Consultant Paediatrician, Sandwell & West Birmingham NHS Trust, UK; hassibnarchi@hotmail.com

doi: 10.1136/adc.2004.057844

At the age of 4 hours, a tachypnoeic, well hydrated, well perfused, and normotensive 34 week preterm neonate develops a metabolic acidosis (arterial pH 7.25, base deficit minus 10 mmol/l). There was no history of perinatal blood loss or sepsis risk factors. No resuscitation was required at birth, and the cord blood pH was 7.3. According to departmental protocol, volume expansion with an intravenous bolus of 10–20 ml/kg of normal saline or 4.5% albumin is advised. You wonder as to the value of this volume expansion.

Structured clinical question

In the absence of asphyxia or hypovolaemia in a newborn infant with metabolic acidosis [patient] does an intravenous bolus of normal saline or albumin [intervention] improve the following [outcomes]: pH, base deficit, mortality, morbidity, length of hospital stay, neurodevelopmental disability?

The outcomes were defined as follows:
- Mortality = neonatal mortality and mortality to discharge.
- Morbidity = peri/intraventricular haemorrhage of any grade, periventricular leukomalacia, patent ductus arteriosus, renal impairment (raised serum creatinine, oliguria), air leak, chronic lung disease (at 28 postnatal days or near term postmenstrual age), necrotising enterocolitis, or retinopathy of prematurity of any grade.
- Neurodevelopmental disability = neurological abnormality including cerebral palsy, developmental delay, or sensory impairment.

Search strategy and outcome

Exclusion criteria: trials including infants with clinically suspected poor perfusion (e.g. low blood pressure, poor cutaneous perfusion) or trials including unselected newborn infants (not known to have metabolic acidosis).

Secondary sources—Cochrane Library (Issue 3, 2003): one protocol—two control trials in CENTRAL, of which one was relevant.

PubMed (1975–2003): search words— (“metabolic acidosis”) AND (“therapy” OR “fluid” OR “albumin” OR “colloid” OR “sodium chloride”) using Clinical Queries methodological filters category (therapy, prognosis) and emphasis (sensitivity, specificity). Limits—Newborn. Search outcome: 87 papers, of which two were relevant (one already retrieved by Cochrane).

SumSearch—96 articles, one protocol (already retrieved by PubMed and Cochrane).

Search results—three articles found, two relevant (already retrieved by PubMed and Cochrane).

Search outcome

See table 2.

Commentary

Previous studies of administering parenteral fluid and/or alkali therapy to neonates with metabolic acidosis have included infants with clinically suspected poor perfusion (e.g. low blood pressure, poor cutaneous perfusion). Other studies of the effect of early volume expansion on mortality and morbidity have included unselected preterm infants not known to have metabolic acidosis.

We found only two studies addressing the benefit of administering intravenous bolus of albumin or normal saline to normovolaemic neonates with metabolic acidosis. They do not however provide a clear answer to the main question of this article in view of few methodological weaknesses. The first study was not blinded. The second study was not randomised and no placebo group was available. Although both studies reported an improvement in the pH and base deficit with volume expansion (although less marked than with bicarbonate), neither of these reports included the following outcomes: survival, morbidity, length of hospital stay, or neurodevelopmental disability. The effect of the resulting correction of the metabolic acidosis on those clinically important outcomes therefore remains unknown.

Table 1 Continued

<table>
<thead>
<tr>
<th>Citation, country</th>
<th>Study group</th>
<th>Study type (level of evidence)</th>
<th>Outcome</th>
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<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alluntas et al (1998), Turkey</td>
<td>47 patients (18 girls, 29 boys; age 4–16 years) of short stature (below third centile for height) without gastrointestinal tract symptoms or endocrinological, cardiac, renal or chromosomal disorders. There were no symptoms associated with coeliac disease or signs of cows’ milk allergy</td>
<td>Cross-sectional study (level 1b/2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>55.3% (26/47) of the short children had biopsy proven coeliac disease</td>
<td>All children were biopsied. Results are not presented clearly; the reader cannot make his own conclusions. Diagnosis of coeliac disease was not according to ESPGAN criteria</td>
</tr>
<tr>
<td>Tumer et al (2001), Turkey</td>
<td>84 children (46 girls, 38 boys; age 16 months – 14 years) with height less than third centile for age; preliminary work-up to evaluate other causes of short stature was found to be negative</td>
<td>Prospective cohort study (level 2b)</td>
<td>Proportion of coeliac disease in the study group</td>
<td>Proportion of coeliac disease was found to be 8.3% (7/84) There is an association between coeliac disease and idiopathic short stature</td>
<td>The IgA EmA test was not verified by a gold standard test (biopsy) in all patients. Diagnosis of coeliac disease was not according to ESPGAN criteria</td>
</tr>
</tbody>
</table>
In the absence of outcome based data, there is currently no evidence to support the routine administration of intravenous bolus of albumin or normal saline to normovolaemic neonates with metabolic acidosis. In addition, as administration of volume expansion was found to be associated with increased mortality in preterm neonates, it is recommended that, in the absence of clear hypovolaemia, caution should be exercised when prescribing volume expansion. A Cochrane protocol reviewing base administration or fluid bolus for preventing morbidity and mortality in preterm infants with metabolic acidosis is currently underway and may provide the appropriate answers.

**Clinical Bottom Line**
- The effect of administration of an intravenous bolus of albumin or normal saline to normovolaemic neonates with metabolic acidosis on mortality, morbidity, and neurodevelopmental outcome in this group of infants is not known.
- There is no evidence to support the benefit of this therapy for such infants.

**References**


**How safe is ibuprofen in febrile asthmatic children?**

**Report by**

A Kader, T Hildebrandt, C Powell, University Hospital of Wales, Cardiff, UK;

**ajmalkader@yahoo.co.uk**

doi: 10.1136/adc.2004.057877

A 4 year old child presents to the paediatric accident and emergency department with a history of fever for 12 hours and clinical signs of an upper respiratory tract infection. The temperature on assessment is 39.5°C. There is a past medical history of asthma. The attending emergency doctor prescribes ibuprofen. The mother is not willing to give ibuprofen to her child, as she was told in the past that it is contraindicated in children with asthma.

**Structured clinical question**

In febrile children with a past medical history of asthma [patient], is ibuprofen in antipyretic doses [intervention], compared to paracetamol [comparison], more likely to cause an acute exacerbation of asthma [outcome]?

**Search strategy and outcome**

Cochrane Databases of Systematic Reviews—none relevant.

Pubmed: “Paracetamol” OR “Ibuprofen” AND “Asthma”.

Limits: All child: 0–18 years, English Language; 30 hits, three relevant.

Journals@Ovid in Athens (ovid) database: “paracetamol” OR “acetaminophen” AND “Ibuprofen” AND “Asthma”; 218 Hits; three relevant (one, same study as above).

See table 3.

**Commentary**

All three references1–3 highlighted by Pubmed using the above search strategy were published by Lesko et al. The data used in all papers were originally derived from the Boston University Fever Study.4 Therefore, the data described in 2002 and 1999 are overlapping.

The original study5 was double blind, randomised, and controlled; patients received either paracetamol in a dose of 12 mg/kg or ibuprofen in a dose of 5 or 10 mg/kg.

To establish the asthma morbidity after the short term use of ibuprofen in children, the data were restricted to include only children being treated for asthma, defined as those who had received a β-agonist, theophylline, or an inhaled steroid on the day before enrolment in the clinical trial. Morbidity from asthma was defined as a report of hospitalisation or outpatient visit for asthma in the month after enrolment.

The data suggested that there was a significantly lower rate of exacerbations of asthma in children receiving ibuprofen compared to children receiving paracetamol. The authors argue that this could be due to the anti-inflammatory action of ibuprofen.

The study by McIntyre and Hull5 was conducted on an inpatient population, which included children with asthma or wheezing. The results showed that no child receiving ibuprofen (including 32 with asthma or a past medical history of asthma) developed symptoms of asthma or wheezing.

In both studies, patients were excluded if there was a known hypersensitivity to paracetamol, ibuprofen, aspirin, or any NSAIDs. Children were also excluded if they had nasal polyps, angioedema, or bronchospastic reactivity to aspirin or other NSAIDs. This small group of children remains potentially vulnerable to ibuprofen or other NSAIDs.

**Table 2** Treatment of metabolic acidosis in normovolaemic newborns

<table>
<thead>
<tr>
<th>Citation</th>
<th>Study group</th>
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<tbody>
<tr>
<td>Dixon et al (1999)</td>
<td>Randomised control trial using 10 ml/kg 4.5% albumin (20 babies) versus sodium bicarbonate (16 neonates) with metabolic acidosis without hypotension</td>
<td>Randomised control trial (level 1b)</td>
<td>Improvement in pH</td>
<td>Change in median pH following bicarbonate (0.10 units) was over twice that in the albumin group (0.04 units)</td>
<td>In normotensive neonates, bicarbonate was more effective than albumin to correct the metabolic acidosis</td>
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</tbody>
</table>
| Belgaumkar et al (1998) | 10 ml/kg 4.5% albumin infusion in 26 ventilated normotensive neonates with metabolic acidosis | Retrospective cohort (level 4) | Improvement in pH and base deficit | Improvement in pH and base deficit up to 6 hours later (p<0.05), no change in blood pressure | *

*Results did not allow risk reduction and NNT to be calculated.

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Arch Dis Child: first published as 10.1136/adc.2004.057844 on 20 August 2004. Downloaded from http://adc.bmj.com/ on September 20, 2023 by guest. Protected by copyright.
Table 3  Ibuprofen in febrile asthmatic children

<table>
<thead>
<tr>
<th>Citation</th>
<th>Study group</th>
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<th>Outcome</th>
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<th>Comments</th>
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<tbody>
<tr>
<td>Lesko et al (2002)²</td>
<td>Total 1879 febrile children 6 mth–12 years receiving asthma medication</td>
<td>Randomised controlled trial</td>
<td>Outpatient visits for asthma or hospitalisation for asthma during 4 weeks post-medication</td>
<td>69 (3.4%) documented outpatient visits: 32 in group 1 and 37 in groups 2 and 3</td>
<td>Ibuprofen was found to be less likely to exacerbate asthma when compared to paracetamol. Children with known hypersensitivity to paracetamol or NSAIDs were excluded</td>
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<td></td>
<td>Group 1: 632 children receiving paracetamol 12 mg/kg/dose</td>
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<td>Group 2: 636 children receiving ibuprofen 5 mg/kg/dose</td>
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<td></td>
<td>Group 3: 611 children receiving ibuprofen 10 mg/kg/dose</td>
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<tr>
<td>McIntyre and Hull (1996)³</td>
<td>Febrile inpatient children (2 mth–12 years)</td>
<td>Randomised controlled trial</td>
<td>Change in axillary temperature, palatability, changes in clinical condition, number and nature of adverse effects</td>
<td>10/76 (13%) patients in the ibuprofen group had 16 adverse events, 14/74 (19%) patients in the paracetamol group had 18 events. This was statistically not significant.</td>
<td>The majority of all adverse events was considered to be either mild or not in relation to the treatment. Children with known hypersensitivity to paracetamol or NSAIDs were excluded</td>
</tr>
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<td>Group 1: 76 received ibuprofen (11 had wheeze or asthma, 21 had a history of asthma or wheezing)</td>
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<td></td>
<td>Group 2: 74 received paracetamol (4 with asthma or wheezing and 12 with a history of asthma or wheezing)</td>
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CLINICAL BOTTOM LINE

- Ibuprofen used as an antipyretic in febrile children with a past medical history of asthma is as least as safe as paracetamol and not likely to exacerbate asthma.
- Ibuprofen should not be used in children with known hypersensitivity to NSAIDs. The possibility of adverse reactions remains in children who have not received NSAIDs at any time in the past.

REFERENCES