

## Contents

	<b>Page</b>
Improving research and access to children's medicines worldwide <i>M Berkovitch, I Choonara, E Jacqz-Aigrain et al.</i>	138
Working with the pharmaceutical industry in relation to paediatric clinical trials. A British perspective <i>O Dewit, R Tiner</i>	140
How long does it take to administer oral medicines to children? <i>C Skwierczynski, S Conroy</i>	145
Paediatric asthma severity score and length of stay in patients presenting to a paediatric emergency department <i>S Chu, J Tan, J A Seabrook et al.</i>	150
Delayed respiratory depression after accidental risperidone overdose <i>H P Satish, H Payne, F Potter et al.</i>	154
Inclusion of a previously published paper	157
Hydroxurea treatment in sickle cell children <i>M-H Odièvre, M Benkerrou, M Belloy et al.</i>	158
A new partnership for drug therapy in children <i>I Choonara, A J Nunn</i>	171
Amniotic fluid disposition of cefazolin during pregnancy <i>K Allegaert, L Lewi, R Verbesselt et al.</i>	172
Paediatric infliximab therapy: patients' and parents' perspectives on treatment options <i>A C M Hazen, F J Smith, K M G Taylor et al.</i>	177
Special thanks	182
Contents for Volume 8	I

## **SHORT COMMUNICATION**

### **Improving research and access to children's medicines worldwide**

Each year 3 million children under the age of five years die from either pneumonia or malaria<sup>1</sup>. The World Health Organization (WHO) recognises and openly states that this situation is unacceptable. The WHO has launched a campaign entitled 'Make medicines child size' which calls for action in relation to both research and access to medicines for children<sup>1</sup>.

WHO points out the urgent need for medicines for neglected diseases such as schistosomiasis, second line treatment of tuberculosis and for co-infection between tuberculosis and HIV. WHO also recognises that children need appropriate formulations and that this needs to include fixed dose combinations for the common infections that affect children in the developing world. These include malaria, tuberculosis and HIV. There is also a need for appropriate antibiotics for neonates with infections.

In its campaign, WHO also emphasises that access to medicines for children throughout the world is crucial. It is unacceptable that children should die from treatable conditions such as pneumonia and malaria. Governments need to work in conjunction with other organisations, including the pharmaceutical industry, to ensure that children who need cheap medicines should be able to access them<sup>2</sup>.

The announcement that WHO will establish a website for clinical trials that have been carried out in children is a major step forward. The existing clinical trial registers do not specifically cater for paediatric studies. The only paediatric clinical trials register is no longer in existence due to lack of support from the major medical journals<sup>3</sup>. This most recent initiative by WHO will be a major step forward for paediatric research.

The European Society for Developmental, Perinatal and Paediatric Pharmacology (ESDP) fully supports this WHO campaign. The ESDP, although predominantly a European organi-

sation, is proud to have members from outside of Europe. We recognise that we have insufficient members in developing countries and are keen to strengthen links with both individuals and institutions in the developing world who wish to improve the treatment of children. This may involve collaborative research or training.

The ESDP recognises that it is the right of children throughout the world to have access to medicines in an appropriate formulation with scientific evidence for both efficacy and safety. This is a major task and cannot be achieved by the ESDP working in isolation. Having seen legislative changes in both North America<sup>4</sup> and Europe<sup>5,6</sup> that should hopefully improve drug therapy for children, the stage is set for the biggest challenge, that of children worldwide.

**Mati Berkovitch**  
*Zerifin, Israel*

**Imti Choonara**  
*Derby, UK*

**Evelyne Jacqz-Aigrain**  
*Paris, France*

**Betty Kalikstad**  
*Oslo, Norway*

**Gerard Pons**  
*Paris, France*

**Anders Rane**  
*Stockholm, Sweden*

**John van den Anker**  
*Rotterdam, Netherlands*

**Bart van Overmeire**  
*Antwerp, Belgium*

**Members of the ESDP Council**

## References

1. [www.who.int/childmedicines/en/index.html](http://www.who.int/childmedicines/en/index.html).
2. Tumwine J K. Equitable access to health care. *BMJ* 2007;335:833-834.
3. Bonati M, Pandolfini C, Rossi V et al. Registering paediatric clinical trials. *Paed Perinat Drug Ther* 2006;7:170-171.
4. Spielberg S P. Paediatric therapeutics in the USA and internationally: an unparalleled opportunity. *Paed Perinat Drug Ther* 2000;4:71-74.
5. Saint-Raymond A, Seigneuret N. Medicines for children: time for Europe to act. *Paed Perinat Drug Ther* 2005;6:142-146.
6. Choonara I, Bonati M. European legislation to improve medicines for children. *Paed Perinat Drug Ther* 2007;8:2-3.

CrossRefs are available in the online published version of this paper:  
<http://www.librapharm.com>  
Paper PPDT-0203\_2, Accepted for publication: 9 January 2008  
Published Online: 13 February 2008  
doi:10.1185/146300908X254189