Choonara, Imti (2011) Safety of new medicines in young children. Archives of Disease in Childhood, 96 (9). pp. 872-873. ISSN 0003-9888

Access from the University of Nottingham repository:

Copyright and reuse:

The Nottingham ePrints service makes this work by researchers of the University of Nottingham available open access under the following conditions.

This article is made available under the Creative Commons Attribution Non-commercial licence and may be reused according to the conditions of the licence. For more details see: http://creativecommons.org/licenses/by-nc/2.5/

A note on versions:

The version presented here may differ from the published version or from the version of record. If you wish to cite this item you are advised to consult the publisher’s version. Please see the repository url above for details on accessing the published version and note that access may require a subscription.

For more information, please contact eprints@nottingham.ac.uk
Safety of new medicines in young children

Inti Choonara

The short report by Tullus (see page 881) highlights safety concerns regarding the use of angiotensin II receptor blockers in preschool children.1 Tullus highlights the deaths of three preschool children out of the 183 children with hypertension who received valsartan or candesartan in two clinical trials. One needs to recognise that deaths do occur in clinical trials. A review of paediatric clinical trials published over a 7-year period identified deaths in 11% of the clinical trials.2 In the majority of cases, it was felt by both the authors of the review and the original authors of the published clinical trials that the deaths were unrelated to the investigational drug. There were, however, six clinical trials terminated early because of significant drug toxicity, as well as two trials where the mortality was higher in the treatment group. Fortunately, the risk of death as a direct result of drug toxicity within paediatric clinical trials appears to be low. It is recognised, however, that drug toxicity is poorly reported in clinical trials.3 A recent review of over 100 published paediatric clinical trials found that less than one in five of the clinical trials reported safety data adequately.3 Clinical trials are not the best way of studying drug toxicity as they are often underpowered to detect severe adverse drug reactions.4 Cases of suspected drug toxicity are often reported as individual case reports. Individual case reports may subsequently be confirmed by retrospective reviews or by a greater understanding of the pathophysiology of the toxicity. A good example of the latter is the cardiovascular collapse of newborn infants following chloramphenicol, which was initially described in three newborn infants in 1959.5 Subsequent studies of the metabolism of chloramphenicol in the neonatal period confirmed a considerably decreased capacity to metabolise chloramphenicol and provided a physiological explanation for the toxicity.6 Retrospective reviews of all the cases of fatal hepatotoxicity secondary to sodium valproate reported in the USA between 1978 and 1984 identified that the risk of hepatotoxicity was greatest in children under the age of 3 years and those receiving other anticonvulsants alongside the valproate.7 The cases of five children who died following the use of a propofol infusion as a sedative in critically ill children were reported in 1992.8 A subsequent review of all the reported cases identified that prolonged duration and high doses were more likely to be associated with development of the propofol infusion syndrome (cardiotoxicity, metabolic acidosis and liver damage) in children.9 Many other suspected cases of drug toxicity, however, are neither confirmed nor refuted. Concern has previously been expressed that paediatric case reports of individual adverse drug reactions are less likely to be reported than adult cases.10

Both the clinical trials of angiotensin II receptor blockers were sponsored by the pharmaceutical industry. This is appropriate as the pharmaceutical company that develops a new product should both sponsor clinical trials and publish the results of the trials. There have unfortunately been several instances where the pharmaceutical company has failed to publish clinical trials when their product has either failed to show efficacy11 12 or been associated with significant toxicity.12-14 A review of industry sponsored clinical trials of antidepressants identified 74 clinical trials registered with the Food and Drug Administration (FDA).11 Thirty-eight of these clinical trials reported a positive benefit of the antidepressant and 37 of these were published. In contrast, 22 of the 36 studies which failed to show efficacy were not published. Additionally, 11 of the 36 studies, which the FDA felt failed to show efficacy, were published with a conclusion suggesting that the drug was effective.

Concern has been expressed by different authors about the interpretation of safety data by the pharmaceutical industry in relation to new products.12-14 Rofecoxib was withdrawn in 2004. A meta-analysis of the risk of cardiotoxicity, however, showed that by the end of 2000, data from over 20 000 patients had confirmed a highly significant greater risk of myocardial infarction.15 The authors felt that rofecoxib should have been withdrawn several years earlier. A review of documents that only became available during legal proceedings against the manufacturer of rofecoxib illustrates how the pharmaceutical company delayed informing the regulatory authorities about an increased number of deaths in the treatment group.16 17 This review also highlights how published articles suggested that the drug was ‘well tolerated’ despite significantly greater deaths in the treatment arm. There have been several new products licensed for use in adults which have had to be withdrawn because of significant drug toxicity. Rimonabant, a treatment for obesity, was withdrawn in 2008 because of its psychiatric toxicity.17 In 2009, the oral β-agonist orciprenaline, a treatment for asthma, was withdrawn because of cardiotoxicity.18 In 2010, rosiglitazone, used for adults with type 2 diabetes, was withdrawn because of the increased cardiovascular risk.19 It is fortunately only a minority of medicines that need to be withdrawn because of drug toxicity. However, one always needs to evaluate the risk of toxicity against the advantages of treatment.

History has also shown that young children may be more prone to specific adverse drug reactions than older children.20 The retrospective review of cases of hepatotoxicity secondary to sodium valproate identified that young children under the age of 3 years were at the greatest risk.9 Antihistamines are widely used in both adults and children. Their use in infants under the age of 1 year, however, has been shown to increase the risk of sleep apnoea, and hence, sudden infant death syndrome.21 One therefore needs to be aware that the safety profile of medicines may be different within different paediatric age groups.

New medicines can result in a significant improvement in reducing morbidity...
and mortality. However, one has also to be aware that the safety profile of a new medicine is not established until it has been extensively used for several years. It would seem sensible to use angiotensin II receptor blockers more extensively in older children, and thus establish more safety data, before rushing to use them in preschool children.

Accepted 14 June 2011
doi:10.1136/archdischild-2011-300450

Competing interests None.
Provenance and peer review Commissioned; internally peer reviewed.

REFERENCES
doi:10.1136/archdischild-2011-300450
Safety of new medicines in young children

Imti Choonara

Arch Dis Child 2011 96: 872-873
doi: 10.1136/archdischild-2011-300450

Updated information and services can be found at:
http://adc.bmj.com/content/96/9/872.full.html

These include:

References
This article cites 21 articles, 5 of which can be accessed free at:
http://adc.bmj.com/content/96/9/872.full.html#ref-list-1

Article cited in:
http://adc.bmj.com/content/96/9/872.full.html#related-urls

Open Access
This paper is freely available online under the BMJ Journals unlocked scheme, see http://adc.bmj.com/info/unlocked.dtl

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/
## Topic Collections

Articles on similar topics can be found in the following collections

- ADC Drug Therapy (80 articles)
- Open access (71 articles)
- Occupational and environmental medicine (94 articles)
- Poisoning (113 articles)
- Child and adolescent psychiatry (paediatrics) (461 articles)
- Unwanted effects / adverse reactions (70 articles)
- Child health (2478 articles)
- Drugs: infectious diseases (610 articles)
- Drugs: psychiatry (50 articles)
- Hypertension (215 articles)
- Medicines regulation (47 articles)
- Clinical trials (epidemiology) (369 articles)
- Epidemiologic studies (1142 articles)
- Metabolic disorders (521 articles)
- Drugs: CNS (not psychiatric) (347 articles)

## Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/