mortality. In Gloucestershire there have been a number of fatal accidents among children related to heavy furniture, blind cords, and diaper bags as well as potentially harmful practices such as cosleeping. Over the last few decades, UK injury prevention programs have halved the number of childhood accidental deaths. There is evidence that community-based campaigns bring about positive behavioral change and can reduce the number of injuries necessitating medical attention. The objective was to explore carer awareness of 4 specific hazards (diaper bags, cord blinds, cosleeping, and heavy furniture) linked to pediatric deaths in the region through the use of questionnaires and a standardized educational poster display.

METHODS: A standardized safety awareness poster board was designed with approved charity leaflets, supported by local council funding. Six hundred poster packs were distributed to public centers across the county. A service evaluation questionnaire was offered to carers or parents of children attending the Children's Center of Gloucestershire Royal Hospital during Child Safety Week. It explored their current safety practices and their thoughts on the usefulness and impact of the poster campaign. The survey was approved by the Trust Research Board and did not need ethical approval.

RESULTS: We obtained 103 questionnaire responses over 5 days, 96% of which were from parents. Almost a quarter of respondents were unaware of accidental deaths relating to diaper bags, although most (82%) kept them out of children's reach. Of the 57 respondents who had cord blinds at home, 26% did not have a safety device attached. Despite prominent national campaigns discouraging cosleeping, 42% of all respondents stated they had slept in the same bed as their children when they were <1 year old. Two-thirds (67%) of respondents reported having secure fixtures in place in their home. Many parents stated they were aware of the hazards highlighted (average 1-10 scale rating, 8.2), and had found the campaign useful (average 1-10 scale rating, 7.3). However, carers perceived the potential to alter current practices as negligible (average 1–10 scale rating, 5.3).

CONCLUSIONS: A poster campaign highlighting hazards implicated in local deaths is deemed useful by parents, but the perceived impact of changing home safety practices is negligible. Additional work through the use of focus groups and parental communication is needed to identify how best to promote safety practices for future campaigns.

URL: www.pediatrics.org/cgi/doi/10.1542/peds.2014-3330BB

Imelda Bennett, Lucy Plumb, Bianca Cuellar Gloucestershire Hospitals NHS Foundation Trust, Gloucestershire, England

# Evaluation of Clinical Features of 238 Cases With Febrile Convulsion

BACKGROUND AND OBJECTIVE: Febrile convulsion (FC) is defined as a seizure triggered by fever in children between 6 months and 5 years of age without an underlying central nervous system infection. It is the most common cause of convulsion in childhood, and 3% to 4% of children experience FC at least once by 7 years of age. The objective was to evaluate clinical features, including demographics, laboratory findings, causes of fever, and FC duration among inpatients diagnosed and treated for FC.

METHODS: A total of 238 patients with the diagnosis of FC between May 2009 and May 2012 were included in the study. Demographic, clinical, and laboratory data of the patients were analyzed.

RESULTS: One hundred thirty-nine patients (58.5%) were male and 99 (41.5%) were female, for a male/female ratio of 1.4. Mean age of patients admitted with a first FC was 2.2  $\pm$  1.1 years. The mean temperature measured rectally during the seizure was 38.7°C  $\pm$  0.5°C. Febrile convulsion was diagnosed as simple type in 198 (83.2%) and complex type in 40 (16.8%) of patients. Thirty-three (13.8%) patients developed a second seizure within 24 hours. Median convulsion duration was 2 minutes (range, 1–5). The most common fever etiology was upper respiratory tract infection, occurring in 131 (55%) cases.

CONCLUSIONS: Benign conditions, such as upper respiratory tract infections, are common causes of FC. A conservative approach is most appropriate in these cases.

URL: www.pediatrics.org/cgi/doi/10.1542/peds.2014-3330CC

Mesut Kocak,<sup>a</sup> Ebru Yýlmaz,<sup>b</sup> Osman Ozdemir,<sup>a</sup> Yusuf Aydýn,<sup>c</sup> Ali Osman Koksal,<sup>a</sup> Deniz Yýlmaz,<sup>a</sup> Aslýhan Araslý Yýlmaz<sup>a</sup> <sup>a</sup>Departments of Pediatrics, and <sup>c</sup>Family Medicine, Kecioren Training and Research Hospital, Ankara, Turkey; and

<sup>b</sup>Department of Family Medicine, Atatürk Training and Research Hospital, Ankara, Turkey

## Trends in Nonpolio Acute Flaccid Paralysis Incidence in India 2000 to 2013

BACKGROUND: Although the incidence of polio acute flaccid paralysis (AFP) has decreased in India, the nonpolio AFP (NPAFP) rate has increased. Nationwide, the NPAFP rate is 11.82 per 100 000 population, whereas the expected rate is 1 to 2 per 100 000 population. We examined the correlates of NPAFP to discern explanations for the increase. The incidence of polio AFP in India has decreased. However, the nonpolio AFP rate has increased since 2000. Follow-up of these cases of nonpolio AFP is not done routinely. However, one-fifth of these cases of nonpolio AFP in the state of Uttar Pradesh (UP) were followed up after 60 days in 2005; 35.2% of patients were found to have residual paralysis, and 8.5% had died. This suggests that the pathology in children being registered as having nonpolio AFP cannot be considered trivial. Therefore, there is a compelling reason to try to determine the underlying causes for the surge in nonpolio paralysis numbers.

METHODS: The data on AFP, polio and nonpolio AFP, and number of polio rounds were examined in each state in each year from 2000 to 2013. Multiple linear regression analysis adjusting for region or state, total and female literacy rate, population density, and per-capita gross domestic product was performed. Differences between states and changes over time were analyzed.

**RESULTS:** NPAFP increased with the number of oral polio vaccine (OPV) doses used (R2 = 25.02%; P < .001). When effect of cumulative doses over the previous years was examined, the NPAFP rate in 2013 best correlated with the cumulative doses received in the previous 7 years (R2 = 57.16%), with 2012 excluded because data for this year were incomplete. This correlation was highly significant (P < .001). On multiple regression analysis, the number of OPV doses was the only factor that showed a positive correlation with the NPAFP rate. The average increase in the NPAFP rate was 1.31 per 100 000 population (*P* < .001; 95% confidence interval, 1.11-1.52) with each dose of OPV. The NPAFP rate in UP and Bihar, which had consistently increased each year until 2011, decreased in the 2 states in 2012, coinciding with a reduction in doses of OPV administered.

CONCLUSIONS: The incidence of NPAFP was strongly associated with the number of OPV doses delivered to the area. A dose–response relationship with cumulative doses over the years was also observed, which strengthens the hypothetical relationship between polio vaccine and NPAFP. The fall in the NPAFP rate in Bihar and UP for the first time in 2012, with a decrease in the number of OPV doses delivered, is evidence of a causative association between OPV doses and the NPAFP rate.

URL: www.pediatrics.org/cgi/doi/10.1542/peds.2014-3330DD

Neetu Vashisht,<sup>a</sup> Jacob Puliyel,<sup>b</sup> Vishnubhatla Sreenivas<sup>c</sup> <sup>a</sup>Jaypee Hospital, Noida, Uttar Pradesh, India; <sup>b</sup>St Stephen's Hospital, Delhi, India; and <sup>c</sup>All India Institute of Medical Sciences, New Delhi, India

#### Algorithm of Risk Group Formation for Having Children With Neural Tube Defects Among Reproductive-Age Women and Differentiated Approach to Spinal Dysraphia Prevention

BACKGROUND AND OBJECTIVE: Congenital neural tube defects (NTDs) are the most common central nervous system congenital defect and among the most common congenital malformations of all organ systems. NTDs are the leading cause of neonatal and infant mortality and childhood disability. The objective was to develop risk groups for having children with NTDs among reproductive-age women based on the detection of endogenous and exogenous risk factors and to offer a differentiated approach to fetal NTD prevention.

METHODS: A retrospective analysis of risk factors in women who gave birth to children with NTD (175 women) and a control group (60 mothers of children without NTD, congenital malformations, or other chromosomal aberrations) and a prospective analysis of the folate cycle metabolic disorders and methylenetetrahydrofolate reductase gene polymorphisms C677T and A1298C were carried out.

**RESULTS:** The inclusion criteria for reproductive-age women in risk groups for fetal NTD should be regarded as the identification of  $\geq 1$  of the following risk factors: history of miscarriages or prenatal fetal death (odds ratio [OR] = 3.4); living in polluted areas and using well water for cooking (OR = 2.7); family history of strokes, heart attacks, varicose veins, thromboembolism, and thrombosis (OR = 3.04); family history of gastrointestinal tract or reproductive system cancer (OR 2.9); family history of congenital malformation (OR 3.9); congenital malformations in other children in the family (OR 4.36); and maternal age >35 years (OR = 2.1). When planning a pregnancy, women from the high-risk group are encouraged to identify levels of homocysteine and folic acid in the blood serum. Revealing hyperhomocystinemia alone or in combination with low folate levels in the blood before conception can be considered predictive of fetal NTD formation. The presence of hyperhomocystinemia is an indication for methylenetetrahydrofolate reductase polymorphism identification to determine preventive measures.

CONCLUSIONS: Forming risk groups for fetal NTD among reproductive-age women and detecting changes in folate metabolism will facilitate preventive measures by determining the timing and amounts of folic acid supplements and dietary recommendations.

URL: www.pediatrics.org/cgi/doi/10.1542/peds.2014-3330EE

Nataly Kotova, Vladyslava Maichuk, Nataly Kononenko Odessa National Medical University, Ukraine

### Trends in Nonpolio Acute Flaccid Paralysis Incidence in India 2000 to 2013 Neetu Vashisht, Jacob Puliyel and Vishnubhatla Sreenivas *Pediatrics* 2015;135;S16 DOI: 10.1542/peds.2014-3330DD

Updated Information & Services	including high resolution figures, can be found at: http://pediatrics.aappublications.org/content/135/Supplement _1/S16.2.full.html
Permissions & Licensing	Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at: http://pediatrics.aappublications.org/site/misc/Permissions.xh tml
Reprints	Information about ordering reprints can be found online: http://pediatrics.aappublications.org/site/misc/reprints.xhtml

PEDIATRICS is the official journal of the American Academy of Pediatrics. A monthly publication, it has been published continuously since 1948. PEDIATRICS is owned, published, and trademarked by the American Academy of Pediatrics, 141 Northwest Point Boulevard, Elk Grove Village, Illinois, 60007. Copyright © 2015 by the American Academy of Pediatrics. All rights reserved. Print ISSN: 0031-4005. Online ISSN: 1098-4275.



Downloaded from pediatrics.aappublications.org by guest on May 5, 2015

# PEDIATRICS®

Trends in Nonpolio Acute Flaccid Paralysis Incidence in India 2000 to 2013 Neetu Vashisht, Jacob Puliyel and Vishnubhatla Sreenivas *Pediatrics* 2015;135;S16 DOI: 10.1542/peds.2014-3330DD

The online version of this article, along with updated information and services, is located on the World Wide Web at: http://pediatrics.aappublications.org/content/135/Supplement\_1/S16.2.full.html

PEDIATRICS is the official journal of the American Academy of Pediatrics. A monthly publication, it has been published continuously since 1948. PEDIATRICS is owned, published, and trademarked by the American Academy of Pediatrics, 141 Northwest Point Boulevard, Elk Grove Village, Illinois, 60007. Copyright © 2015 by the American Academy of Pediatrics. All rights reserved. Print ISSN: 0031-4005. Online ISSN: 1098-4275.



Downloaded from pediatrics.aappublications.org by guest on May 5, 2015