

current guidelines for safety during flight for infants with a history of neonatal lung disease.

Authors' affiliations

K Udomittipong, P D Sly, Clinical Sciences, Telethon Institute for Child Health Research and Centre for Child Health Research, University of Western Australia, Perth, Australia
S M Stick, G L Hall, School of Paediatrics and Child Health, University of Western Australia, Perth, Australia
S M Stick, M Verheggen, J Oostryck, P D Sly, G L Hall, Respiratory Medicine, Princess Margaret Hospital, Perth, Australia

KU was funded by the Siriraj Hospital, Thailand. PDS and SS are funded by the National Health and Medical Research Council, Australia.

Competing interests: none.

REFERENCES

- 1 **Samuels MP**. The effects of flight and altitude. *Arch Dis Child* 2004;**89**:448–55.
- 2 **Speizer C**, Rennie CJ 3rd, Breton H. relevance of in-flight medical emergencies on commercial airlines. *Ann Emerg Med* 1989;**18**:26–9.
- 3 **Coker RK**, Partridge MR. Assessing the risk of hypoxia in flight: the need for more rational guidelines. *Eur Respir J* 2000;**15**:128–30.
- 4 **Lee AP**, Yamamoto LG, Relles NL. Commercial airline travel decreases oxygen saturation in children. *Pediatr Emerg Care* 2002;**18**:78–80.
- 5 **Barry PW**, Pollard AJ. Altitude illness. *BMJ* 2003;**326**:915–9.
- 6 **Carpenter TC**, Niermeyer S, Durmowicz AG. Altitude-related illness in children. *Curr Probl Pediatr* 1998;**28**:177–98.
- 7 **British Thoracic Society**. Managing passengers with respiratory disease planning air travel: British Thoracic Society recommendations. *Thorax* 2002;**57**:289–304.
- 8 **Oades PJ**, Buchdahl RM, Bush A. Prediction of hypoxaemia at high altitude in children with cystic fibrosis. *BMJ* 1994;**308**:15–8.
- 9 **Buchdahl RM**, Babiker A, Bush A, *et al*. Predicting hypoxaemia during flights in children with cystic fibrosis. *Thorax* 2001;**56**:877–9.
- 10 **Parkins KJ**, Poets CF, O'Brien LM, *et al*. Effect of exposure to 15% oxygen on breathing patterns and oxygen saturation in infants: interventional study. *BMJ* 1998;**316**:887–91.
- 11 **Buchdahl R**, Bush A, Ward S, *et al*. Pre-flight hypoxic challenge in infants and young children with respiratory disease. *Thorax* 2004;**59**:1000.
- 12 **Jobe AH**, Bancalari E. Bronchopulmonary dysplasia. *Am J Respir Crit Care Med* 2001;**163**:1723–9.
- 13 **Poets CF**, Samuels MP, Southall DP. Potential role of intrapulmonary shunting in the genesis of hypoxemic episodes in infants and young children. *Pediatrics* 1992;**90**:385–91.

LUNG ALERT

Reliability of a pulmonary embolism management algorithm

▲ van Belle A, Buller HR, Huisman MV, *et al* for the Christopher Study Investigators. Effectiveness of managing suspected pulmonary embolism using an algorithm combining clinical probability, D-dimer testing, and computed tomography. *JAMA* 2006;**295**:172–9

The combination of clinical scoring systems and D-dimer testing is increasingly being used to stratify probabilities of pulmonary embolism (PE). This study was targeted to assess the clinical effectiveness of an algorithm using clinical decision rules, D-dimer testing, and CT scans in patients with suspected PE. The primary outcome measures were fatal or symptomatic venous thromboembolism (VTE) at 3 months follow up.

A total of 3306 eligible patients with suspected PE (age >18 years, 57.4% female, 18.3% inpatients) were prospectively enrolled from 12 hospitals. Using a modification of Well's clinical decision rules, PE was considered unlikely in 2206 (66.7%). PE was excluded in 1057 (32%) of those following a normal D-dimer test (1149 had an abnormal test), but 29 of the 1057 were anticoagulated for other reasons. At 3 months, five patients (0.5%) suffered VTE. Clinically, 1100 (33.3%) were considered likely to have had a PE.

For the above 2249 (1100+1149) requiring a CT scan, PE was excluded in 1505 (69 were anticoagulated for other reasons, 18 (1.3%) suffered VTE), CT scanning confirmed PE in 674 patients, 20 CT scans were inconclusive (one VTE), and 50 did not have a CT scan (two VTE). The prevalence of PE was 12.1% (226/2206) in the clinically "unlikely" group compared with 37.1% (408/1100) in the clinically "likely" group ($p < 0.001$).

The authors conclude that this diagnostic strategy is effective and associated with a low risk of subsequent VTE. Limitations of this study include the use of two different kinds of CT scanner (multidetector and single detector) with potentially different pick up rates. The prevalence of PE of 23.2% (266/1149) in the "clinically unlikely PE and abnormal D-dimer" group shows that a clinical scoring system alone is inadequate.

S Bari

Specialist Registrar in Respiratory Medicine, Cardiothoracic Centre and Royal Liverpool University Hospitals, Liverpool, UK; baris@doctors.org.uk