CASE REPORT

A case of neonatal stridor

V Abdullah, S K Ng, S N Chow, F T Yau, C A van Hasselt

A 13 day old baby girl presented with severe inspiratory stridor and respiratory distress. She was the first child of the family born spontaneously with a birth weight of 3290 g. The antenatal history was uneventful. There was no premature rupture of the membranes and the liquor was clear. The baby received no airway instrumentation other than routine nasal suction with a plastic catheter immediately after delivery. Noisy breathing was present from day 2 of life. On day 5, the noise increased and sucking in of the chest was noticed. The condition then further deteriorated, and she was admitted to hospital on day 13. She was tachypnoeic (respiratory rate of 60/min) and stridorous with severe subcostal recession. Pulse oximetry measured 88–92% at 3 litres/min oxygen through the nasal cannula. She had no fever and her heart rate was 200/minute. Her white cell count was 27.8 × 10⁹/l (neutrophils 48.2%), blood gases were pH 7.36, PaO₂ = 4.81 kPa, PaCO₂ = 10.40 kPa, HCO₃⁻ = 39.9 mmol/l. Intravenous fluid was given, and antibiotics, including ampicillin, netilmicin, and metronidazole, were started on a 14 day course of antibiotics, and she made an uneventful recovery. The mother’s high vaginal swab was taken for culture, which also grew GBS.

DISCUSSION

Retropharyngeal abscesses commonly present in infants and young children. In the series of Coulthard and Isaacs, neonates only accounted for 10% of the cases. The dangers of a retropharyngeal abscess are airway obstruction and spread of infection to involve the carotid sheath and/or the mediastinum. Prompt treatment is therefore indicated. The airway, if compromised, should be secured before other diagnostic procedures; this is especially important in neonates. Oral intubation should be carried out with great caution to avoid rupturing the abscess, which could result in aspiration of its contents with fatal consequences. Parenteral antibiotics covering both aerobic and anaerobic flora of the upper aerodigestive tract should be prescribed. A CT scan is mandatory to establish the diagnosis and rule out other rare congenital retropharyngeal masses, such as cystic hygroma, haemangioma, and retropharyngeal goitre. These exclusions are essential in a neonate because they may not, as illustrated by this case, present with typical signs of sepsis. A CT scan is also useful in delineating the extent of the disease, which helps in planning surgical approaches. A retropharyngeal abscess with a cavity medial to the great vessels is best drained transorally, whereas those abscesses that extend laterally to the parapharyngeal space should be approached by a lateral cervical incision with the placement of surgical drains.

This is the first report of GBS infection presenting as neonatal stridor as a result of a retropharyngeal abscess. In addition to the well described early and late onset GBS infection, it seems possible for the organism to localise in the retropharyngeal space. The maternal genitourinary tract is the usual location of the organism, which was probably the source of the early infection in this child. We can only speculate on how the organism becomes inoculated and localised in the retropharyngeal space. Universal antenatal screening for GBS is not practised in Hong Kong. Most gynaecologists would screen for GBS in women who are symptomatic, and give antibiotics accordingly. Intrapartum chemoprophylaxis is definitely indicated for subsequent deliveries in this mother. GBS infection is one of the most common causes of neonatal sepsis, and it has presented uniquely in this case as a retropharyngeal abscess.
Retropharyngeal abscess caused by group B streptococcus

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