Fatal neonatal septicaemia and meningitis due to *Haemophilus influenzae* acquired from the mother

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**Summary**

A fatal case of neonatal *Haemophilus influenzae* type b, septicaemia and meningitis is described. Although the mother was asymptomatic, maternal genital tract culture yielded *H. influenzae* which was indistinguishable from the strain found in the infant. Despite the rarity of this organism as a cause of neonatal sepsis, increased awareness is essential so that this important offending pathogen is not missed.

KEY WORDS: neonatal, *Haemophilus influenzae*, septicaemia, meningitis, maternal genital tract.

**Introduction**

*H. influenzae* is rarely grown from the female genital tract (Hurley, 1970) and in a recent survey of cervical cultures among normal and pregnant women the isolation rate was less than in 1% (Khuri-Bulos and McIntosh, 1975). Though a rare cause of neonatal infection, it is one of the commonest pathogens in children under the age of 3 years (Fothergill and Wright, 1933). As most adults possess serum bactericidal antibody against *H. influenzae*, the scarcity of neonatal infections has been attributed to the passive transfer of protective antibody to the baby. Therefore, any well documented rise in incidence of *H. influenzae* type b during the past few years may be attributed to absence or low level of antibody against this pathogen. However, this has not proved to be the case and the age distribution of *H. influenzae* meningitis has not changed during the past four decades (Peter, 1973).

**Case report**

A 2780 g boy was born to a 31-year-old primigravida. Gestational age was estimated to be 37 weeks. Rupture of the membranes occurred 22 hr prior to an assisted breech delivery. At birth the infant had an Apgar score of 4 at 1 min and 9 at 5 min and the condition was fair. The only method of resuscitation was to clear the airways by mucus extractor. Forty-eight hours after birth the baby suddenly collapsed and was transferred to the Special Care Baby Unit where he began to have convulsions.

Cerebrospinal fluid (CSF) examination revealed red cells 1.48 x 10^9/1 white cells 0.54 x 10^9/1 (all polymorphonuclear leucocytes), Gram negative rods ++ +, which on culture grew *H. influenzae* type b, sensitive to ampicillin and chloramphenicol. CSF protein was 1.65 g/l and glucose 1.1 mmol/l. Blood cultures also grew the same organism. No pathogens were isolated from nose, throat, ear, umbilical, rectal swabs, end of endotracheal tube and from pharyngeal suction. The white cell count was 3.3 x 10^9/1 and the serum calcium 1.69 mmol/l.

Treatment was started with ampicillin and chloramphenicol (300 mg and 35 mg intravenously 4 times a day respectively) with phenobarbital and dexamethasone. At this stage throat and high vaginal swabs were collected from the mother were which latter on culture yielded *H. influenzae*. The three isolates of *H. influenzae*—from the CSF and blood of the infant and the vagina of the mother were identified to be type b, confirming that the baby acquired it from the mother (Dr D. C. Turk, Public Health Laboratory, Northern General Hospital, Sheffield). Despite ventilation and other supportive measures, in view of the baby’s rapidly deteriorating condition with convulsions, cyanosis, grey colour, apnoea and laboured respiration, a second lumbar puncture was performed 72 hr after the first one. This showed Gram negative rods but was sterile on culture. However, his condition deteriorated further with right upper lobe pneumonia, pulmonary haemorrhage, evidence of necrotizing enterocolitis, and severe neurological changes; he died on the 7th day.

**Discussion**

Like Group B streptococcal infection, the pathogenesis of neonatal *H. influenzae* infection appears...
likely to be due to transmission of organisms from
tooth to baby at or immediately before delivery.
That infection can occur before birth is highlighted
by the fact that there is a report of septic abortion
infected with *H. influenzae* (Berczy, Fernlund and
Kamme, 1973), and high incidence of prematurity
and bacteraemia in infants with early onset *H.
influenzae* sepsis (Lillien et al., 1978). Contrary to
what was anticipated, genital carriage rate is very
low (White and Koontz, 1968; Khuri-Bulos and
McIntosh, 1975). Nevertheless it was invariably
present in the genital tract of mothers whose babies
developed sepsis (Khuri-Bulos and McIntosh, 1975;
Granoff and Nankervis, 1975; Nicholls, Yuille and
Mitchell, 1975; Marston and Wald, 1976; Bale and
Watkins, 1978). Unfortunately, except those men-
tioned above, in other reports there was either failure
to collect any swabs or to record results of culture
reports from mothers. In this particular case the
mother was asymptomatic with no vaginal discharge
or pyrexia and there was no relevant immediate past
history to suggest carriage of such pathogens.

Although this baby was not premature, he suc-
cumbed not only because of overwhelming infection
due to septicemia and meningitis, but also because of
very low white count, low calcium, and sudden
collapse at 72 hr which needed intubation leading to
pulmonary haemorrhage terminally. This is only the
second report of neonatal *H. influenzae* sepsis in the
United Kingdom where the same organism has been
demonstrated to be present in the maternal genital
tract.

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