Chickenpox pneumonia: an association with pregnancy

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ABSTRACT Three pregnant patients with chickenpox pneumonia are described. A review of reports published since 1965 together with these cases shows that 28 of the 46 women (61%) were pregnant, and 21 were in the last trimester. Although the disease is more common and more severe in pregnancy, the proportion of deaths was no greater than usual. In this retrospective analysis there was no evidence that treatment with acyclovir conferred benefit but this may have reflected its use only in very ill patients.

Introduction

In our hospital over two years we have seen three pregnant women with chickenpox pneumonia. This prompted a search of reports published worldwide to find whether chickenpox pneumonia was associated with pregnancy and whether pregnancy affected the outcome.

Case reports

CASE 1
A 27 year old woman at 16 weeks' gestation was admitted with a five day history of rash and dyspnoea, of two days' duration. She was centrally cyanosed and dyspnoeic at rest, and had a widespread erythematous maculopapular rash with vesicles and harsh breath sounds throughout both lung fields.

Chest radiography showed extensive confluent mottling throughout both lung fields. Arterial blood gas analysis while she was breathing air showed that pH was 7-42, oxygen tension (PaO2) 2-3 kPa, and carbon dioxide tension (PaCO2) 4-3 kPa. She was initially treated with 90% oxygen, which caused the PaO2 to rise to 6-9 kPa while the PaCO2 remained unchanged. As the patient was exhausted, however, ventilation was instituted and nine days later a tracheostomy was performed. Her condition improved subsequently, the tracheostomy was closed, and the patient was discharged 23 days after admission. Blood gas tensions and the chest radiograph had returned to normal. She delivered a healthy female infant by spontaneous vaginal delivery at 40 weeks' gestation.

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appearance. Microscopy showed bacterial superinfection but no specific changes attributable to chickenpox pneumonia.

CASE 3
A 25 year old woman at 36 weeks' gestation presented with a two day history of increasing dyspnoea and dry cough. Four days before this she had developed a chickenpox rash. At presentation she was febrile (38°C), with a typical chickenpox rash, a respiratory rate of 25/min, and tachycardia (130 beats/min). Examination of the chest showed nothing remarkable.

The chest radiograph showed bilateral confluent mottling. Arterial blood gas analysis on admission showed pH to be 7·38, PaCO₉ 3·5 kPa, and PaO₂ 5·5 kPa. The patient was given 60% oxygen and intravenous acyclovir 10 mg/kg eight hourly. Her condition deteriorated over the next six hours and an emergency caesarean section was performed. Assisted ventilation was continued postoperatively. A healthy baby girl was delivered, who developed chickenpox six days after birth. The mother was ventilated for a total of five days and had acyclovir for seven days. Recovery was complicated by a deep venous thrombosis in the right calf. Mother and baby were discharged three weeks later. Arterial blood gases and the chest radiograph had returned to normal.

Review of previous reports

A review of the reports published since 1965¹–³⁴ combined with our three cases yields a description of 99 adults with chickenpox pneumonia (table 1).

Of the 46 women (in nine others the sex was not stated), 28 were pregnant, indicating a clear association with pregnancy. The disease occurred in the third trimester in 21 and in the second in three; the duration of pregnancy was not stated in four. As judged by the number requiring assisted ventilation, chickenpox pneumonia appeared to be a more serious illness in pregnancy, but mortality (11%) was similar to that in non-pregnant patients. Treatment with antiviral agents did not appear to confer benefit; five of 44 patients (11%) receiving acyclovir and none of six patients receiving other antiviral agents died, compared with five of 49 (10%) patients who received no antiviral drug (χ² = 0·178, p > 0·5 for summation of results). The benefit from acyclovir could be obscured if patients were given the drug only if they were more ill than those who were not. When adult mortality in the era before the introduction of acyclovir was compared with that in the period after its introduction no significant difference was found (table 2).

When the mother survived infant mortality was low and only one infant died. Six infants were delivered during the illness and 15 after it. Only two infants born alive developed chickenpox.

Discussion

The varicella-zoster virus is a highly contagious DNA virus. Five per cent of women in the child bearing age lack specific IgG and are therefore susceptible. Varicella has been thought to be relatively infrequent in pregnancy, with an incidence of 0·7 per 1000 pregnancies.²³ In the United States the mean incidence for women of 15–45 years of age is 2·2/1000/year.

Carstairs and Emond²⁶ in 1963 reported 120 patients from published reports with acute chickenpox pneumonia, and Knvyett (1965)²⁷ described a personal series of 35 cases. In these earlier descriptions clinical data were scanty and Knvyett did not record the age or sex incidence, or whether any of the patients were pregnant. For these reasons we have confined our analysis to patients described since 1965.

Only four of the cases that occurred during pregnancy occurred outside the last trimester. This may be explained by the progressive changes in the immune system that occur in pregnancy,²⁸ which is the immune system

Table 2  Numbers of patients surviving before and after the introduction of acyclovir

<table>
<thead>
<tr>
<th></th>
<th>Survived</th>
<th>Died</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before acyclovir</td>
<td>29</td>
<td>1</td>
</tr>
<tr>
<td>Acyclovir era</td>
<td>60</td>
<td>9</td>
</tr>
</tbody>
</table>

χ² = 1·23, p > 0·02.

Table 1  Adult chickenpox pneumonia: a review of previously published cases combined with the present three (number (%))

<table>
<thead>
<tr>
<th>Sex</th>
<th>Ventilated</th>
<th>Died</th>
<th>Acyclovir used</th>
<th>Other antiviral agents†</th>
<th>Fetus died</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non-pregnant (n = 71)</td>
<td>M 44</td>
<td>13 (18)</td>
<td>28 (39)</td>
<td>2 (3)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>F 18</td>
<td>(718†)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pregnant (n = 28)*</td>
<td></td>
<td>13 (46)</td>
<td>3 (11)</td>
<td>16 (57)</td>
<td>4 (14)</td>
</tr>
</tbody>
</table>

*Twenty one in last trimester and four unknown.
†Not stated.
‡Cytosine arabinoside, idoxuridine, gammaglobulin, adenine arabinoside.
being required to mount a normal immune response to exogenous antigen while not mounting a response against the foreign antigen of the fetal trophoblast. This task is accomplished by several changes in the immune system: the thymus becomes progressively involuted owing to increased concentrations of oestradiol and corticosteroids, the number of monocytes is increased (though there are conflicting reports about the number of T and B cells), and there is an increase in leucocyte phagocytic and bactericidal responses. Other abnormalities, such as antibody and allergen responses, are less clearly defined. These immunological disturbances of pregnancy may be mediated by pregnancy associated globulins. Pregnancy is associated with an increased incidence of influenza, cytomegalovirus and *Haemophilus influenza* infections, and tuberculosis. Early delivery of the infant during the illness is unnecessary to save the mother’s life. This is at variance with the experience in Lassa fever.

Only two infants developed chickenpox after delivery. In both cases delivery occurred within four days of the mother’s developing chickenpox, allowing no time for her protective IgG to cross the placenta to protect the infant.

Antiviral treatment, in particular the introduction of acyclovir in 1979, does not appear to have altered the outcome and its role in treatment is unproved.

Diffuse pulmonary calcification may occur three to 10 years after chickenpox pneumonia, and this was present in three of 21 patients who underwent prolonged follow up. In the present review of 99 patients none had pulmonary calcification, but this probably reflects the brief duration of follow up.

Patient 3 was under the care of Dr J Davies and we are grateful for his permission to describe the case.

References


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