Delayed presentation of vernix caseosa peritonitis

AC Chambers, AV Patil, R Alves, JC Hopkins, J Armstrong, RN Lawrence

Great Western Hospitals NHS Foundation Trust, UK

ABSTRACT

INTRODUCTION Vernix caseosa peritonitis (VCP) is a rare and poorly recognised condition resulting from a sustained foreign body reaction to the vernix caseosa of the baby. This case-based review aims to highlight its importance for any medical team managing patients with peritonitis who have undergone a recent Caesarean section.

CASE REPORT A 31-year-old woman presented 5 weeks after a Caesarean section with symptoms and signs of peritonitis.

CONCLUSIONS Laparotomy and peritoneal lavage is the mainstay of treatment for VCP. Knowledge of the condition may stop inadvertent resection of normal intra-abdominal organs. Greater awareness of VCP is required to ensure earlier recognition as patients can recover well following timely operative intervention.

KEYWORDS Vernix caseosa peritonitis – Caesarean section – Laparotomy

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CORRESPONDENCE TO
Adam Chambers, Department of General Surgery, Great Western Hospital, Marlborough Road, Swindon SN3 6BB, UK
T: +44 (0)1793 604 020; E: adam.chambers@nhs.net

Vernix caseosa is a thick white paste that is present on foetal skin from the third trimester of pregnancy until shortly after birth.1 It is unique to our species and consists predominantly of water (80%) combined with proteins and lipids in equal portions (20%). Biologically, vernix caseosa is beneficial to the neonate and infant via production of antimicrobial peptides2 as a moisturiser for neonatal skin and providing a waterproof barrier allowing for maturation of the neonatal stratum corneum. It is thought to be the cause of vernix caseosa peritonitis (VCP).3

VCP is a rare condition that affects women post-Caesarean section following spillage of amniotic fluid and vernix caseosa into the peritoneal cavity. Vernix caseosa can result in a profound systemic inflammatory response that necessitates maternal laparotomy and may lead to erroneous resection of intra-abdominal organs. Diagnosis is often difficult due to a lack of awareness of the condition and may only be made following histological examination. Confusion regarding diagnosis can result in considerable anxiety for both the patient and surgeon in the ongoing management of the condition.

We present a case report of VCP in a 30-year-old woman 5 weeks after a Caesarean section.

Case history

A 30-year-old primigravida woman delivered a healthy baby boy by lower segment Caesarean section (LSCS) at 40 weeks after conversion from natural delivery due to failure to progress. She had an unremarkable antenatal period and no significant medical history. Five weeks following the LSCS, she presented with a 48-hour history of generalised abdominal pain, vomiting and diarrhoea, and was admitted with a provisional diagnosis of gastroenteritis under the care of the medical team. Clinical examination revealed a systemic inflammatory response syndrome (SIRS) with a pulse rate of 150bpm, blood pressure of 97/55mmHg and temperature of 39.1ºC. Abdominal examination demonstrated a soft abdomen with generalised tenderness. The LSCS wound looked healthy. The C-reactive protein level was 481mg/l with a neutropaenia of 2.2 x 10⁹/l.

Intravenous fluid and antibiotics were administered. Computed tomography (CT) of the abdomen and pelvis on day 2 demonstrated some free fluid. The patient was transferred to the intensive care unit (ICU) and was prescribed granulocyte colony stimulating factor, which increased her neutrophil count to 41.2 x 10⁹/l. Stool, urine and blood cultures were all negative.

The patient deteriorated further with increased pain, a distended abdomen and signs of peritonitis. Repeat CT demonstrated an increased volume of intra-abdominal free fluid. An urgent laparotomy was performed, which revealed a multiloculated fluid collection in the abdomen and pelvis that was not malodorous. The uterus was intact. There was widespread inflammatory change throughout the peritoneal cavity with large volumes of fibrinous material. Appendicitis was presumed and an appendicectomy and extensive peritoneal lavage performed. Two large drains were sited, in the right paracolic gutter and the pelvis. The patient improved post-operatively with a concomitant decrease in inflammatory markers.

Histology of the appendix revealed acute fibrinous sepsis and material taken from the abdominal cavity dem-
DElAYED PRESENTATION Of VERNix CASEOSA PERITONITIS

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onstrated degenerate anucleate squamous cells in keeping with VCP (Fig 1). Further cultures taken from the drain post-operatively revealed no evidence of intra-abdominal infection. The patient was discharged pain free 25 days after surgery.

Discussion

The literature base for VCP is limited (Table 1) and, to date, only 20 cases of VCP following a Caesarean section have been published.3–15 It is unclear whether two of the cases of VCP described by Herz et al5 had been published previously. The largest series of three patients is from Australia.16 Most of the cases, however, were from the US.17 VCP has also been seen antenatally following spontaneous rupture of membranes. Caesarean sections performed on two women revealed VCP at the time of delivery, presumably as a result of reflux of vernix caseosa into the peritoneal cavity.18

Our patient presented to hospital five weeks following her LSCS, which is the longest period currently described in the literature. Diagnosis was made after seven days, multiple radiographic tests, a period on the ICU, and following laparotomy and histological confirmation. This highlights the diagnostic dilemma with VCP.

A study published in 2011 reports that VCP could be diagnosed with image guided biopsy15 although this may not be practical in the presence of peritonitis and SIRS. Most women in this cohort of patients, in corroboration with our case, progressed to emergency laparotomy and organ resection for presumed diagnoses of more common causes of peritonitis (appendicitis, bowel/ureteric injury or uterine rupture). Our experience suggests such difficulty in diagnosis can result in considerable psychological stress for the patient.

Following our experience with VCP, we recommend laparotomy and peritoneal lavage as the mainstay of treatment, with intravenous antibiotic cover and critical care support if required. We note that steroid therapy has been used successfully in some cases.11 However, in our case we felt it was not appropriate. Prognosis appears to be good with the majority of patients recovering following surgery. Nevertheless, some authors report recurrent vernix caseosa abscess formation necessitating drainage or reoperation.12,16

Although the condition is not often described in medical textbooks,17 with the increasing number of Caesarean sections performed worldwide, this diagnosis must be considered in post-Caesarean section women presenting with abdominal pain and signs of peritonitis.

Conclusions

This case highlights the lack of awareness in the medical community of VCP. In spite of its rarity, VCP represents serious post-Caesarean section morbidity and may require urgent surgical management. Diagnosis requires close involvement with the histopathologist. Management should entail rigorous peritoneal lavage and placement of intra-abdominal drains to monitor for early recurrence. It is unknown whether VCP can present later than five weeks post-partum but consideration of VCP as a diagnosis is important.

Figure 1  Histology of appendix and intra-abdominal tissue:
The pan-cytokeratin stain demonstrates a mesothelial reaction on the serosal surface of the appendix and over the serosal exudates (A; 10x magnification). At higher power one can see both nucleated cell staining with cytokeratin (mesothelial cells) and anucleate linear structures consistent with anucleate squamous cells (B; 200x magnification). Both nucleated mesothelial cells and anucleate cells consistent with squamous cells are seen in the exudates from the peritoneal cavity (C; 400x magnification).
## Table 1: Case reports of vernix caseosa peritonitis following Caesarean section

<table>
<thead>
<tr>
<th>Authors</th>
<th>Cases</th>
<th>Caesarean to acute presentation</th>
<th>Operation performed</th>
<th>Resected organs</th>
<th>Histopathology</th>
<th>Microbiology</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Krumerman and Pouliot, 1976</td>
<td>1</td>
<td>28 days</td>
<td>Laparotomy</td>
<td>Appendix</td>
<td>Appendiceal serositis, giant cells surrounding anucleate squames</td>
<td>Peritoneal material culture -ve</td>
<td>Recovered</td>
</tr>
<tr>
<td>Herz et al., 1982</td>
<td>2</td>
<td>a) 8 days; b) 14 days</td>
<td>a) Laparotomy + adhesiolysis; b) Laparotomy</td>
<td>a) None; b) Hysterectomy + BSO</td>
<td>a) Granulomatous reaction + squames, lanugo hair, keratin and meconium; b) Tubo-ovarian abscess + squames engulfed by foreign body giant cells</td>
<td>a) N/A; b) Exudate cultured B melaninogenicus</td>
<td>Recovered</td>
</tr>
<tr>
<td>Freedman et al., 1983</td>
<td>2</td>
<td>a) 9 days; b) 17 days</td>
<td>a) Laparotomy; b) Laparotomy</td>
<td>a) Appendix; b) Hysterectomy + BSO</td>
<td>a) Foreign body reaction + keratin debris and hair fragments; b) Tubo-ovarian abscess + squames surrounded by giant cells</td>
<td>N/A</td>
<td>N/A</td>
</tr>
<tr>
<td>Boothby et al., 1985</td>
<td>1</td>
<td>35 days</td>
<td>Laparotomy</td>
<td>Descending colon</td>
<td>Colon not perforated, giant cells, macrophages with degenerating superficial squamous cells</td>
<td>N/A</td>
<td>Recovered</td>
</tr>
<tr>
<td>George et al., 1995</td>
<td>2</td>
<td>a) 8 days; b) 10 days</td>
<td>a) Laparotomy; b) Laparotomy</td>
<td>a) Ascending colon; b) Omentum</td>
<td>a) Serositis, granulomatous inflammation with anucleate squamous cells, lanugo hair; b) mixed inflammation with anucleate squamous cells</td>
<td>a) Peritoneal fluid culture -ve; b) Culture -ve</td>
<td>Recovered</td>
</tr>
<tr>
<td>Zellers and Balaj, 1996</td>
<td>2</td>
<td>a) 12 days; b) 5 days</td>
<td>a) Laparotomy; b) Laparotomy</td>
<td>a) Appendix; b) Appendix</td>
<td>Both a) and b): Periappendicitis + foreign body reaction with anucleate squamous cells</td>
<td>N/A</td>
<td>Recovered</td>
</tr>
<tr>
<td>Nuñez, 1996</td>
<td>1</td>
<td>3 days</td>
<td>Laparotomy</td>
<td>Ascending colon</td>
<td>Not perforated, peritonitis, foreign body granulomas with anucleated squames</td>
<td>N/A</td>
<td>Recovered</td>
</tr>
<tr>
<td>Mahmoud et al., 1997</td>
<td>1</td>
<td>3 days</td>
<td>Laparotomy</td>
<td>Appendix</td>
<td>Anucleated squamous cells</td>
<td>N/A</td>
<td>Recovered with steroid therapy</td>
</tr>
<tr>
<td>Tawfik et al., 1998</td>
<td>1</td>
<td>4 days</td>
<td>Laparotomy</td>
<td>None</td>
<td>Mixed inflammatory exudate + squames, hair shafts and meconium pigment</td>
<td>Blood + aspirate cultures -ve</td>
<td>Further abscess drained</td>
</tr>
<tr>
<td>Cummings et al., 2001</td>
<td>1</td>
<td>6 days</td>
<td>Laparoscopy</td>
<td>Gallbladder</td>
<td>Granulatation tissue aggregated about anucleate squames; no acute cholecystitis</td>
<td>Gram -ve</td>
<td>Recovered</td>
</tr>
<tr>
<td>SelO-Ojeme et al., 2007</td>
<td>1</td>
<td>7 days</td>
<td>Laparotomy</td>
<td>None</td>
<td>N/A</td>
<td>Scanty growth of Gram positive</td>
<td>Recovered</td>
</tr>
<tr>
<td>Stuart et al., 2009</td>
<td>3</td>
<td>a) 5 days; b) 8 days; c) 4 days</td>
<td>a) Laparotomy; b) Laparotomy; c) Laparoscopic to open + 2nd laparotomy</td>
<td>a) Omental biopsy; b) None; c) Appendix at 2nd operation</td>
<td>a) Serositis, granulomas + fetal squames; b) Chronic granulomas + fetal squames; c) Acute inflammation surrounding fetal squames and keratin debris from 1st operation</td>
<td>a) N/A; b) Urine culture -ve; c) Urine + blood cultures -ve</td>
<td>a) Recurrent vernix caseosa abscesses; b) 7 months’ pain; c) Recurrent obstruction</td>
</tr>
<tr>
<td>Wisanto et al., 2010</td>
<td>1</td>
<td>24 days</td>
<td>Laparoscopy</td>
<td>None</td>
<td>Anucleated squamous cells</td>
<td>N/A</td>
<td>Recovered</td>
</tr>
<tr>
<td>Myers and Fernado, 2011</td>
<td>1</td>
<td>21 days</td>
<td>Fine needle aspiration</td>
<td>None</td>
<td>Mixed inflammatory response with anucleate and mature squamous cells</td>
<td>Culture -ve</td>
<td>Recovered with no operation</td>
</tr>
</tbody>
</table>

-ve = negative; BSO = bilateral salpingo-oophorectomy; N/A = not applicable
in any woman presenting with an acute abdomen following a recent Caesarean section.

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References