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Toxic megacolon during pregnancy in ulcerative colitis: A case report

Ayyaz Quddus^{a,*}, Beatriz Martin-Perez^b, Henry Schoonyoung^b, Matthew Albert^b, Sam Atallah^b

^a Department of Gastroenterology, Royal Free Hospital, London, United Kingdom ^b Florida Hospital Division of Colon and Rectal Surgery,Orlando, FL, USA

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ABSTRACT

INTRODUCTION: Ulcerative colitis is an idiopathic inflammatory bowel condition whose peak incidence coincides with fertility in female patients. In pregnancy, acute fulminant colitis is rare, and, when it becomes refractory to maximum medical therapy, emergency colectomy is mandated. Over the past quarter century, there have been few reports of this rare event in the literature.

PRESENTATION OF CASE: We report a 26 year old primigravida female who presented with toxic megacolon during the third trimester of pregnancy, unresponsive to medical therapy. She subsequently underwent an urgent low transverse caesarean section with a total colectomy. Both mother and child made a satisfactory recovery post operatively.

DISCUSSION: Although the fetus is at higher risk than the mother in such a circumstance, morbidity and mortality rates are still noticeably high for both, and therefore, prompt diagnosis is key.

CONCLUSION: It is imperative that female patients planning to conceive with a known diagnosis of ulcerative colitis liaise with their obstetricians and gastroenterologists early to optimise medical treatment to prevent the development of a toxic megacolon and that conception is planned during a state of remission. Should surgical intervention become required, this can be performed with favourable outcomes for mother and child, as demonstrated in this report.

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1. Introduction

Ulcerative colitis (UC) was first described by the British physician Sir Samuels Wilks in 1859 [1]. It is an idiopathic inflammatory bowel condition, depicted as a chronic, relapsing disease whose intestinal manifestations are restricted to the colon [1,2], although a subset of patients may also develop ileal inflammation or so-called 'backwash ileitis' [2,3]. It has been estimated that between 30 and 50% of female patients with UC will have an exacerbation during pregnancy or in the early postpartum period [4–7]. Patients with active disease at the time of conception may have an increased risk of spontaneous abortion. Rates of preterm delivery, low birth weight (<2500 g), and congenital anomalies are also increased [8]. A rare but potentially fatal complication of UC is toxic megacolon. This condition is characterised pathologically by severe, transmural inflammation with infiltration of the colonic wall by neutrophils, lymphocytes, histiocytes and plasma cells [9]. The most significant risk of toxic megacolon is perforation into the surrounding peritoneal cavity which holds a considerable mortality rate due to the sequalae of sepsis [10-12]. The development of toxic megacolon

E-mail address: ayyazquddus@googlemail.com (A. Quddus).

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during pregnancy is an unusual occurrence. Only seven authors with a combined eleven cases have been described in the literature (Table 1) [11–17]. Here, we illustrate a case report of a 26 year old female that developed toxic megacolon at 31 weeks gestation.

2. Case report

A 26 year old primigravida female with an eight year history of ulcerative colitis presented to the emergency department with abdominal pain, fever, bloody diarrhoea, and generalised weakness. At this initial presentation, the patient was at 26 weeks gestation and had poorly controlled colitis requiring tri-modal therapy with mesalazine, 6-mercaptopurine and prednisolone. The patient was stabilised initially with conservative management including analgesia, intravenous fluids and intravenous steroids and was subsequently discharged. She was then readmitted at 31 weeks gestation with systemic signs of sepsis, including tachycardia, hypotension and a pyrexia of 38.3 °C. Physical examination demonstrated a distended abdomen consistent with pregnancy, which was exquisitely tender on palpation and percussion with positive rebound tenderness indicative of peritonitis. Plain film abdominal X-ray demonstrated severe colonic distension without free air (Fig. 1). The patient underwent an emergent caesarean section via an infra-umbilical midline incision. The fetus was successfully

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^{*} Corresponding author. Present address: Royal Free Hospital, Pond Street, London NW3 2QG, United Kingdom. Tel: +44 20 7794 0500.

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84 **Table 1**

Summary of reported cases of toxic megacolon in pregnancy.

Authors	Year	n	Maternal gestation (weeks) ^a	Maternal mortality (%)	Fetal mortality (%)	Pooled fetal and maternal mortality (%)
Marshak et al. [15]	1960	1	24	0	100	50
Peskin and Davis [16]	1960	1	24	0	100	50
Holzbach [17]	1969	2	22,30	50	100	75
Becker [12]	1972	1	23	0	0	0
Cooksey et al. [13]	1985	1	24	0	0	0
Anderson et al. [14]	1987	3	33,34,37	0	33	17
Ooi et al. [11]	1985	2	10,16	0	0	0
Total	-	11	-	-	-	-

^a Represents the gestational period at which the diagnosis of toxic megacolon was established.



Fig. 1. Preoperative plain film abdominal radiograph of a 26 year old lady with suspected toxic colitis. The single frontal supine view demonstrates a diffusely distended transverse colon with displacement by the gravid uterus.

delivered with apgar (appearance, pulse, grimace, activity, respiration) scores of 9 at one minute and 9 at five minutes; weight 1.474 kg and height 36.2 cm. Immediately after delivery, the midline incision was extended approximately 2 cm cephalad and a total abdominal colectomy with end ileostomy was performed. Total operative time for the combined procedures was 96 min and estimated blood loss was 250 ml. The patient was discharged nine days postoperatively in stable condition; while the neonate was stabilised in the intensive care unit, where, ventilator support was initially required. He was weaned off of the ventilator, transitioned to a neonatal progressive care unit and eventually discharged home two weeks after birth. Pathologic examination of the surgical specimen confirmed the diagnosis of toxic megacolon with transmural inflammation and erosion of the bowel wall in regions (Fig. 2). Four weeks post-operatively, the patient underwent completion proctectomy and creation of an ileal reservoir with construction of an ileoanal anastomosis and diverting ileostomy. The reason for accelerating the procedure was based on patient choice and the fact that she had recovered very well from her emergency colectomy. Operative time was 177 min and estimated blood loss was 200 ml. The loop ileostomy was reversed three months later (Fig. 3).

Two months following the loop ileostomy reversal the patient was seen in clinic reporting 2–3 bowel movements a day,

constitutionally well with optimal healing of her surgical wounds. The effect of reduced fertility due to the ileal pouch-anal anastomosis (IPAA) formation in the future was explained during the consultation. The neonate measured between 40 and 50th centile in all his growth assessments and he also made a satisfactory recovery from the procedure.

3. Discussion

Although ulcerative colitis has a peak incidence which coincides with females of childbearing age, it does not affect fertility in women unless they have undergone curative treatment in the form of a proctocolectomy plus IPAA [18]. Several meta-analyses have found that IPAA conferred a two to three fold increased risk of infertility compared with medical management [19,20] The most likely mechanisms for impaired fertility following surgery are higher rates of hydrosalpinx, destruction of fimbria and tubal obstruction due to adhesions and scarring [21].

Studies have shown that disease status at the time of conception plays a vital role in predicting disease activity during pregnancy and also on the outcome of the pregnancy [22]. When conception occurs during a period of remission in inflammatory bowel disease then the risk of relapse during pregnancy is one-third and these patients are more likely to have quiescent disease activity during pregnancy [23]. This is the same risk of relapse in nonpregnant patients with Crohn's disease. Conversely, women with UC or Crohn's have a twofold risk of developing exacerbations during pregnancy if during conception they had active disease [24,25]. The underlying cause for the increased risk is not clearly understood.

In addition, active inflammatory bowel disease at the time of conception and during pregnancy has been shown to be associated with a high risk pre-term birth, low birth weight and small for gestational age birth compared with women with IBD whose disease is in remission during conception and during the course of the pregnancy [26–28]. Other predictors of adverse pregnancy outcomes include family history of IBD, disease localisation and IBD surgery [26–28].

Females that have an exacerbation of ulcerative colitis during their pregnancy (most likely first trimester [13]) or early postpartum period, are initially treated medically. Ulcerative colitis during pregnancy that is refractory to medical therapy can lead to the development of toxic megacolon. Over the past quarter of a century, there have been no reports of this rare event. This is perhaps because of improvements in the medications used to treat inflammatory bowel disease, including new classes of medicines – most notably recombinant monoclonal antibodies which target TNF-alpha (e.g. Infliximab). Previous literature demonstrates that different surgical options have been applied to the pregnant patient with a toxic megacolon [11-17]. Despite these variances is surgical approach, the objective is twofold: (a) save the mother by removing the diseased organ and (b) deliver the fetus, if at a viable gestation age. It is clear that fetal mortality is higher than maternal mortality, Download English Version:

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