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Postpartum Bowel Perforation after Normal Vaginal Delivery without History of Inflammatory Bowel Disease

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Background	Ischemia complicates up to 42 percent of bowel perforations and increases the mortality in these patients significantly. Bowel perforation is a very rare occurrence in any post-partum patient, but it can be seen more commonly following a Caesarean section (C-section) than following a normal vaginal delivery (NVD). It is more commonly described in postpartum patients who have a history of chronic bowel diseases or prior abdominal surgery, but this is still a very rare phenomenon.
Summary	This case presents a previously healthy, 22-year-old female with no surgical history who developed a perforation of the right mid-colon five days post-normal vaginal delivery (NVD). At this time, she presented to the emergency department (ED) with severe, progressive abdominal pain, signs of sepsis and multiple organ failure. Imaging revealed pneumoperitoneum. The working differential included: Ogilvie syndrome, Clostridium difficile (C. diff) colitis, volvulus and a possible ischemic event. She was taken emergently to the operating room for an exploratory laparotomy. Obstetrics and gynecology (OBGYN) was present during this intervention and noted the uterus was normal for postpartum day five, ruling out uterine cause of the pneumoperitoneum noted on imaging. A right hemicolectomy with anastomosis and a diverting loop ileostomy was performed at this time. Postoperatively, the patient recovered in the intensive care unit and one week later she had abdominal fluid collections that could not be drained by interventional radiology and required an additional exploratory laparotomy for adequate drainage. Ten days post-second laparotomy she was discharged home. Her ileostomy was reversed two months after discharge without issue.
Conclusion	A colonic perforation as the cause of our patient's abdominal pain was very low on the differential for this young, healthy woman with no past medical or surgical history, however we can see from this case that it can happen. Therefore, abnormal abdominal distension and pain in the post-partum period should at least warrant an abdominal X ray.
Keywords	Bowel obstruction, peritonitis, postpartum, ischemic bowel changes, sepsis, pneumoperitoneum, right hemicolectomy, diverting loop ileostomy, fecal peritonitis, vaginal delivery, Ogilvie syndrome

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Case Description

A previously healthy, 22-year-old African American female, with no surgical history, G1P1001 had a normal vaginal delivery (NVD) induced at 41 weeks. Her peripartum period was complicated by preeclampsia with severe features that required magnesium before delivery through 24 hours postpartum and a positive group B streptococcus culture. Postpartum day one she complained of some constipation and abdominal pain. There were no peritoneal signs on physical exam. Abdominal X ray showed mildly distended loops of small bowel with a normal amount of fecal material in the right hemicolon. She was given two enemas and had a bowel movement (BM) prior to discharge. She and her infant were discharged three days postpartum; of note in the discharge summary her abdomen was mildly distended and tender to palpation, with no peritoneal signs, these findings were deemed normal for the postpartum period. She was sent home with the following bowel regimen: MiraLAX®, Phillips'® Milk of Magnesia as needed, and encouraged ambulation—also a normal occurrence for postpartum patients.

On postpartum day five, the patient returned to the ED with signs of sepsis and multiple organ failure. She was complaining of severe abdominal pain that had been progressively worsening for five days, nausea and vomiting, subjective fever, chills, sore throat, dyspnea, chest pain, palpitations, weakness, diffuse muscle cramps and lower leg edema. Her last BM was on the day of readmission, it was liquid and unclear if it was bloody because of postpartum vaginal bleeding. Her labs were unremarkable except for a leukocytosis of nineteen and renal failure with a BUN of 54 and creatinine of 2.7. On examination, the patient appeared very fatigued and drowsy, tachycardia was noted with a regular rhythm. The abdomen was grossly distended, diffusely tender with rebound and guarding. There were hypoactive bowel sounds and tympany present. The rest of the physical exam was unremarkable. Abdominal CT scan showed pneumoperitoneum.



Figure 1. CT scan of abdomen done in ED on admission.



Figure 2. CT scan of abdomen done in ED on admission, lung window.

The differential diagnoses included: Ogilvie syndrome, volvulus, C. diff colitis or possibly a postpartum ischemic event. Surgery consulted OBGYN due to the unusual presentation so close to the delivery and performed an emergent exploratory laparotomy; the uterus was normal size for postpartum day five ruling out a uterine cause, such as infection, uterine perforation, etc. for the pneumoperitoneum. Upon further exploration, a perforated right colon, close to the hepatic flexure with fecal peritonitis was found. Due to the significant inflammation and peritonitis present, a clear etiology was not evident intra-operatively. Because of the patient's instability, a quick but thorough inspection and palpation of the distal colon was done. There was no colonic redundancy to suggest volvulus. There were no gross tumors, any inflammation, or distension to suggest obstruction or infection. The rest of colon was seemingly very normal. A right hemicolectomy with anastomosis and diverting loop ileostomy was performed and two drains were left. The pathology workup of the specimen showed no abnormalities. One week post-operatively the patient had some fluid collections that could not be drained by interventional radiology and required an open washout with multiple drains placed. She recovered well and was subsequently discharged after ten days.

Her diverting loop ileostomy was reversed three months after discharge. The reversal was performed after thorough evaluation, including a barium enema and colonoscopy without any abnormal findings.

Discussion

A literature review over a three month period using many sources including but not limited to: DynaMed, New England Journal of Medicine, UpToDate, PubMed and Google scholar, revealed only three similar published cases of post-NVD bowel perforation; the latest of which was published in June 2014. All three aforementioned case reports cite Ogilvie syndrome as the cause of bowel perforation post-NVD.1, 2, 3 Ogilvie syndrome is a functional obstruction that can eventually progress to perforation, this can rarely present after a surgical procedure, such as a C-section. This presentation, though rare, is much more common than post-NVD. One published article describes post-NVD bowel perforation in a patient who had a significant history of symptomatic ulcerative colitis, including a previous proctocolectomy and ileo-pouch-anal anastomosis.4 The second case described an older patient who was having her second child via vaginal delivery.2 The third case is also an older woman post-NVD and her symptoms presented immediately after delivery of the fetus³. Based on the aforementioned research, this is the first case report of its kind.

Our differential diagnoses for this patient included, *C. diff* colitis because of the antibiotic use before discharge due to a suspected urinary tract infection. A fecal leukocyte count was unremarkable, and stool *C. diff* PCR was negative. Volvulus was deemed unlikely during the surgery because the colon was not distended and no signs of volvulus were noted intraoperatively. Because of these findings, our working diagnosis was Ogilvie syndrome, pseudo obstruction complicated by ischemia and finally leading to right colon perforation. Ischemia complicates up to forty-two percent of all bowel obstructions (including patients not in the peripartum period) and significantly increases mortality and morbidity.¹

The postpartum period can rarely lead to colonic pseudo obstruction of unclear etiology. This pseudo obstruction can lead to colonic dilation, ischemia of the bowel wall and eventually perforation. This disease process is very rarely found post-NVD. The etiology of Ogilvie syndrome is still unknown, but it has been known to occur with abdominal trauma, sepsis and abdominal or pelvic surgery. Making the diagnosis of Ogilvie syndrome is difficult because the symptoms are very nonspecific. The most common symptom noted in patients with Ogilvie syndrome is abdominal distension. One article reported that the association between Ogilvie syndrome and vaginal delivery maybe due to the declining serum estrogen levels in the postpartum period.

Ogilvie syndrome that leads to perforation usually involves ischemia of the bowel wall. Other ischemia-related conditions can occur in the peripartum period due to a state of hypercoagulability during pregnancy. The most common ischemic event during the peripartum period is venous thromboembolism (VTE), this can occur anywhere, but commonly in the mesenteric, splenic, portal, pelvic and deep veins of the leg. Other postpartum ischemia-related conditions are: Sheehan syndrome, hypovolemic acute renal failure, ischemic heart disease and ischemic stroke. VTE in the postpartum period has a prevalence of 1 in 1600. Maternal deep vein thrombosis is more often associated with PE than as an isolated finding, PE is the seventh leading cause of maternal mortality (nine percent).

The clinical signs of abdominal distension in the postpartum period should warrant a high suspicion for further evaluation despite these patients being generally healthy.

Conclusion

Colonic perforation is extremely rare in postpartum patients, but can happen, even in otherwise healthy women. The initial symptoms of abdominal distension and pain are vague, and sometimes expected in this population, but should raise consideration for further work-up or monitoring as the results can be devastating.

Lessons Learned

Postpartum patients with any abnormal abdominal distension or tenderness should be given a thorough evaluation. At the very least, these patients should have an abdominal x-ray performed and follow up if symptoms do not resolve clinically. This early diagnostic consideration will reduce morbidity and mortality for patients similar to the one presented in this case.

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