Case Report

Diaphragmatic Hernia (Bochdalek Hernia) in an Adult: a Case Report

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Abstract:

Bochdalek hernia (BH) is the most common type of congenital diaphragmatic hernia and constitutes 85% of cases. In adults BH is extremely rare. Most of them are asymptomatic. We present a case of BH in an adult patient to point out the importance of clinical examination and computed tomography (CT) scan in reaching the exact diagnosis. A 17-year-old male was presented to the surgery department with a 3-days history of recurrent abdominal pain, vomiting and unable to lie flat. The BH was confirmed on computed tomography (CT) imaging and the patient underwent emergency laparotomy and per-operatively we found part of stomach, greater omentum, splenic flexure of colon along with spleen herniated into the left hemithorax through a defect of approximately 8x6cm in the left postero-lateral aspect of the diaphragm. Reduction of hernial contents and surgical repair of the defect was done with non-absorbable suture. His post-operative period was uneventful. Although rare, this disorder should be recognized and treated appropriately to avoid complications.

Key words: Bochdalek hernia, Laparotomy.

Introduction:

Diaphragmatic hernia occurs in a defect of the diaphragm, through which loops of small and large bowel, stomach, liver, and spleen may protrude into the thoracic cavity of the involved side. This defect of the diaphragm can be either congenital or acquired after a blunt trauma¹². The incidence of congenital diaphragmatic hernias (CDH) is 1 in 2500 births and left sided CDH is more common than right-side (85% and12%, respectively)³. Although CDH is diagnosed prenatally or in the immediate postnatal period, in 5–25% of cases diagnosis can be late, and could be detected during routine examinations or examination because of respiratory or gastrointestinal problems³. Bochdalek hernia (BH) is a variety of congenital diaphragmatic hernia caused by failure fusion of the posterolateral diaphragmatic foramina; resulting in the displacement of abdominal contents into thoracic cavity⁴. Bochdalek first described this anomaly in 1848⁴. BH is very rare beyond childhood with an incidence of 0.17%. Diaphragmatic hernias are usually asymptomatic in adults⁵. In 5–10% of affected individuals, signs and symptoms of appear later in life and may include respiratory or GI problems⁶.

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Symptomatic hernias are misdiagnosed in up to 38% of cases leading to unnecessary interventions such as chest tube insertion or failure to diagnose which could lead to strangulation of the herniated content and death⁷.
Case report:

A 17-year-old boy came to our surgery department with the complaints of recurrent upper abdominal pain for last 1 year which was burning in nature, insidious on onset, non-radiating and aggravated after taking meal. The pattern of pain changed to continuous and agonizing nature with increasing severity for last 3 days before hospital admission with vomiting for several times for last 24 hours. He gave no history of surgery or trauma. On examination, abdomen was diffusely tender, maximum over epigastric & left hypochondriac region, with rigidity and muscle guarding. Chest movement was normal, vocal fremitus was reduced on left side, percussion note was dull and on auscultation breath sound was diminished on left lower chest with presence of bowel sounds in the left hemithorax. Plain X-ray abdomen showed loops of bowel were high-up, lying in the left lower hemithorax. There was no evidence of pneumoperitoneum or abnormal air fluid level (Figure-1).

CT scan of chest showed large bowel loop along with peritoneal fat entered into left hemithorax through abnormal opening in the postero-lateral aspect of diaphragm, left lung is collapsed & mediastinum was shifted to the right (Figure-2).

Figure 1: Plain X-ray abdomen showed loops of bowel were high-up, lying in the left lower hemithorax, no evidence of pneumoperitoneum or abnormal air fluid level was seen.

Figure 2: Coronal section of CT scan of chest showing large bowel loop along with peritoneal fat entered into left hemithorax.

Other laboratory investigations were within normal limit. At first, the patient was resuscitated. Then an emergency laparotomy was arranged. After opening the abdomen, we found part of stomach, omentum, splenic flexure of colon along with spleen herniated into the left hemithorax through a defect of approximately 8x6cm cm in the postero-lateral aspect of the diaphragm. Reduction of hernial contents was done meticulously, viability checked and primary repair of the defect was done by 1-0 polypropylene suture (Figure 3).

Figure 3: Showing the defect of diaphragm
Abdominal drain tube was kept in the site of repair & a chest drain tube was kept for 3 days. He was kept nothing per oral for 3 days. Post-operative period was uneventful (Figure-4).

Figure 4: Post-operative chest x-ray P/A view showing chest drain tube in situ, normal contour of left dome of diaphragm, gastric bubble below the left dome of diaphragm.

Discussion:
Congenital diaphragmatic hernias, clinically represented in adult life are extremely rare, and they can occur through the frontal parasternal opening (Morgagni hernia 1-6%) or through the postero-lateral opening (Bochdalek hernia 80-90%)8. Postero-lateral (Bochdalek hernia) is the most common type of congenital diaphragmatic hernias that occurs in 95% of cases, resulting from inadequate closure of the poster lateral pleuropitoneal membranes9. The overall prevalence of Bochdalek hernias in adults is 6-10%10. Literature review reveals that defects occur more frequently on the left side (80-90%) than on the right side of the diaphragm, and the hernia contents included omentum, left colon, stomach, small bowel loops and spleen11. In adults, most Bochdalek’s hernias are asymptomatic, and thus the finding of the condition is incidental. Some patients have no symptoms and the disorder is unexpectedly detected on a chest X-ray or CT scan of the thorax. Symptomatic cases of Bochdalek’s hernia in adults are rare. They are diagnosed in adults as an incidental medical finding or when the symptoms appear. Our patient presented with abdominal pain, vomiting and unable to lie flat. Acute presentation is usually due to incarceration, obstruction, or strangulation of the hernia. Diagnosis is confirmed by a combination of chest X-ray and computed tomography (CT). Chest CT is usually the imaging of choice12.

The management of Bochdalek hernia includes reduction of hernial contents to the peritoneal cavity and repair of the diaphragmatic defect13. Various surgical repair options include open surgery, laparoscopic repair, and thoracoscopic approach. Although laparotomy was the most widely used surgical approach, minimally invasive surgical techniques have gained popularity since their first report in 199514. Laparoscopic repair can be performed with a low complication rate (7%) and short hospital stay14. Several authors suggested an abdominal approach for left-sided defects and a thoracic approach for right-sided hernias11. Regardless of the type of surgical procedure, suturing the defect is likely important for the restoration of the anatomy between the thoracic and abdominal cavities. In patients with large diaphragmatic hernias, synthetic or biologic grafts can be used for definitive treatment and prevention of recurrence but it has some complication like adhesion formation with diaphragm15. In our patient the size of the defect in the diaphragm was 8 cm wide. We decided to open repair of the hernia instead of attempting laparoscopic repair. Nevertheless, a tensionless type of repair has been validated as an option for BHs, which is similar to the type of repair used for all other hernia repairs14. Diaphragm hernia repair approaches can cause postoperative pneumonia, deep venous thromboembolism, myocardial infarction, or sepsis15. In our case, none of these complications occurred.

Conclusions:
Patients with a Bochdalek hernia may not have any symptom and the condition may present as an acute abdomen or with respiratory problems. Examination of the chest may give strong suspicion and chest radiography is a good screening tool but CT scan has higher sensitivity for this lesions. Even though rare, this disorder should be recognized and treated appropriately to avoid complications.

References:


