

Unveiling the Chilaiditi Syndrome: A Case Report and Management Implications

Review began 03/24/2024

Review ended 03/28/2024

Published 04/02/2024

© Copyright 2024

Kamel et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 4.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Ibrahim Kamel ¹, Yusuf Yalcin ², Reid Ponder ², Ibrahim Elkhawas ³, Zeeshan Solangi ¹

1. Internal Medicine, Steward Carney Hospital, Boston, USA 2. Radiology, Tufts University School of Medicine, Boston, USA 3. Pulmonary Critical Care, Baystate Medical Center, Springfield, USA

Corresponding author: Zeeshan Solangi, zeeshan.solangi@steward.org

Abstract

The Chilaiditi syndrome is when the radiologic Chilaiditi sign, defined by the interpositioning of the colon between the liver and diaphragm, becomes complicated by clinical symptoms such as respiratory insufficiency or bowel obstruction. We present the case of a 70-year-old male with a history of depression, anxiety, gastroesophageal reflux disease (GERD), and post-polio syndrome, who presented with left shoulder pain, chronic weakness, and dizziness. Initial evaluation revealed hypotension and elevated lactic acid, attributed to dehydration. Further imaging identified a Chilaiditi sign, thus raising suspicion of small bowel obstruction and the Chilaiditi syndrome. Despite conservative management, the patient continued to experience elevated lactic acid levels, prompting a computed tomography (CT) angiogram to rule out bowel ischemia. No acute intra-abdominal pathology was identified, and the patient improved with hydration and bowel rest. This case highlights the challenges in diagnosing and managing the Chilaiditi syndrome in the setting of chronic comorbidities.

Categories: Gastroenterology, Internal Medicine, Radiology

Keywords: lactic acidosis, computed tomography, colonic interposition, radiologic finding, chilaiditi syndrome

Introduction

The Chilaiditi sign is a rare, incidental radiologic finding secondary to colonic interposition between the liver and diaphragm. The Chilaiditi syndrome then arises if a patient develops associated clinical symptoms. Current literature reports a sign incidence between 0.025% and 0.28% on plain film, although computed tomography (CT) evaluation is more sensitive with incidence estimates between 1.2% and 2.4% [1-3]. The etiology of the Chilaiditi sign involves both congenital and acquired factors, ranging from ligament abnormalities, right diaphragm paralysis, chronic constipation, redundant colon, liver cirrhosis, and multiple pregnancies [4]. The mechanism is poorly understood, although it is thought to depend on both a hypermobile intestine and an enlarged subdiaphragmatic space [5].

As the Chilaiditi sign and syndrome are characterized by apparent subdiaphragmatic air, several dangerous conditions can present similarly on imaging. This list includes pneumoperitoneum, subphrenic abscess, diaphragmatic hernia, and, in the context of trauma, diaphragmatic rupture [6-8]. Management for these etiologies is surgical and frequently urgent. In contrast, the Chilaiditi syndrome can be a self-resolving or chronic entity and could therefore be managed with a more conservative approach [9]. We present a case of a 70-year-old male who presented with left shoulder pain, chronic weakness, and dizziness and was found to have hypotension, elevated lactic acid, and a Chilaiditi sign. We discuss our clinical reasoning and management of this patient.

Case Presentation

A 70-year-old male with a history of depression, anxiety, gastroesophageal reflux disease (GERD), and post-polio syndrome presented to our emergency department with left shoulder pain, chronic weakness, and dizziness. He was found to be hypotensive by a visiting nurse and brought to the emergency department. Initial evaluation at our center revealed hypotension (87/61 mmHg), elevated lactic acid (4.6 mmol/L), and normal oxygen saturation. The chest X-ray shown in Figure 1 was unremarkable for acute cardiopulmonary pathology but was remarkable for subdiaphragmatic air.

How to cite this article

Kamel I, Yalcin Y, Ponder R, et al. (April 02, 2024) Unveiling the Chilaiditi Syndrome: A Case Report and Management Implications. Cureus 16(4): e57483. DOI 10.7759/cureus.57483

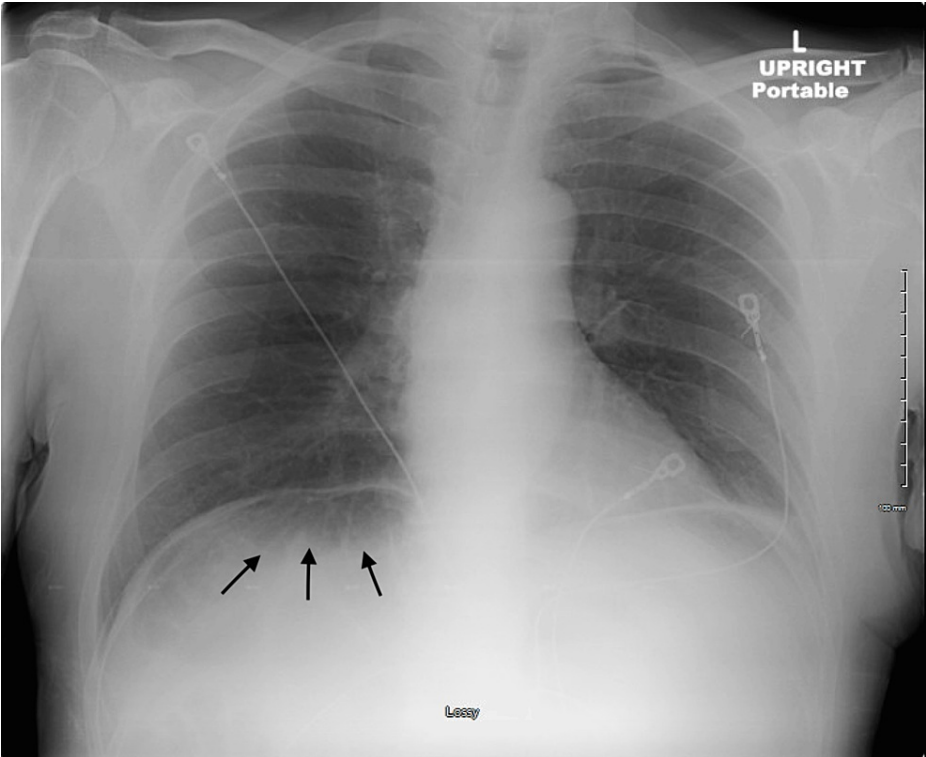


FIGURE 1: Chest X-ray

Chest X-ray showing bowel loops under the diaphragm. The black arrows pointing at the bowel loops

Further review demonstrated overlying bowel loops, suggesting that this air was intraluminal and secondary to an interposed intestine. The patient’s chart was reviewed for the anatomical variant of the Chilaiditi sign, and a CT of the abdomen from one year prior showed haustral markings between the abdominal wall and liver, consistent with the Chilaiditi sign. The results of this study are shown in [Figure 2](#).

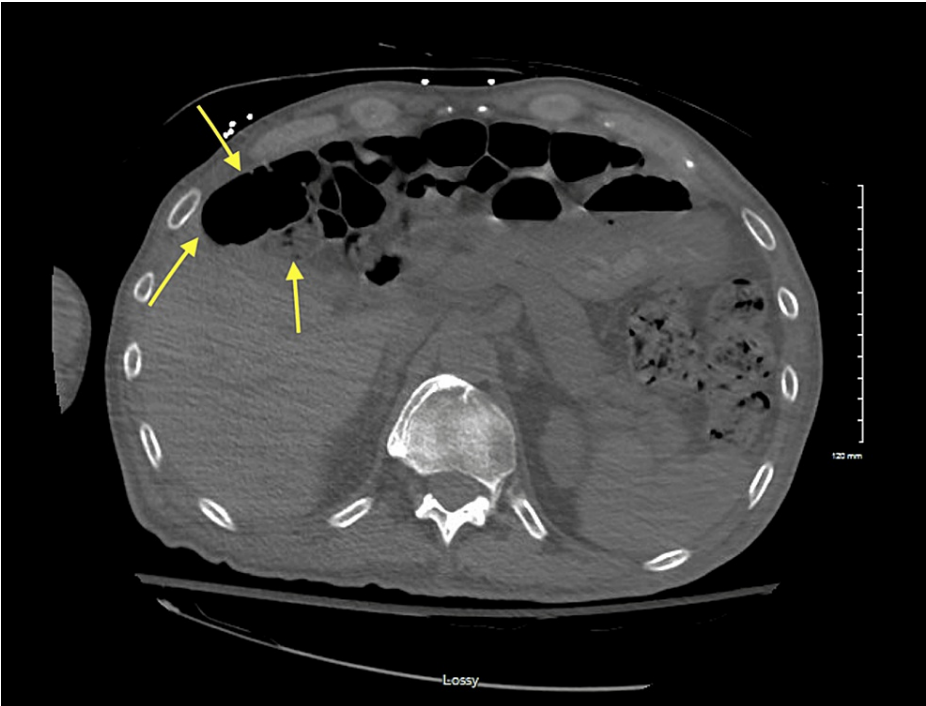


FIGURE 2: Abdominal CT one year prior to presentation

Image showing CT scan of the abdomen with the bowel loops between the abdominal wall and the liver. The arrows pointing at the bowel loops

CT: computed tomography

Further evaluation showed no clinical signs of infection, abdomen that was soft, and no tenderness or distention. The patient improved with IV hydration. However, lactic acid levels continued to increase, prompting a CT of the abdomen and pelvis shown in Figure 3, which again revealed the Chilaiditi sign with suspicion of small bowel obstruction.



FIGURE 3: Abdomen/pelvis

CT of the abdomen/pelvis with white arrows pointing at the bowel loops interposed between the abdominal wall and the liver, which are dramatically changed from the previous study

CT: computed tomography

Notably, this study contained more interposed bowel loops compared to the CT from one year prior. Surgery recommended bowel rest and close monitoring, especially given the patient's deconditioning and overall frailty.

Despite bowel rest and hydration, lactic acid levels fluctuated, necessitating a CT angiogram to rule out bowel ischemia. The results of this test are shown in Figure 4.

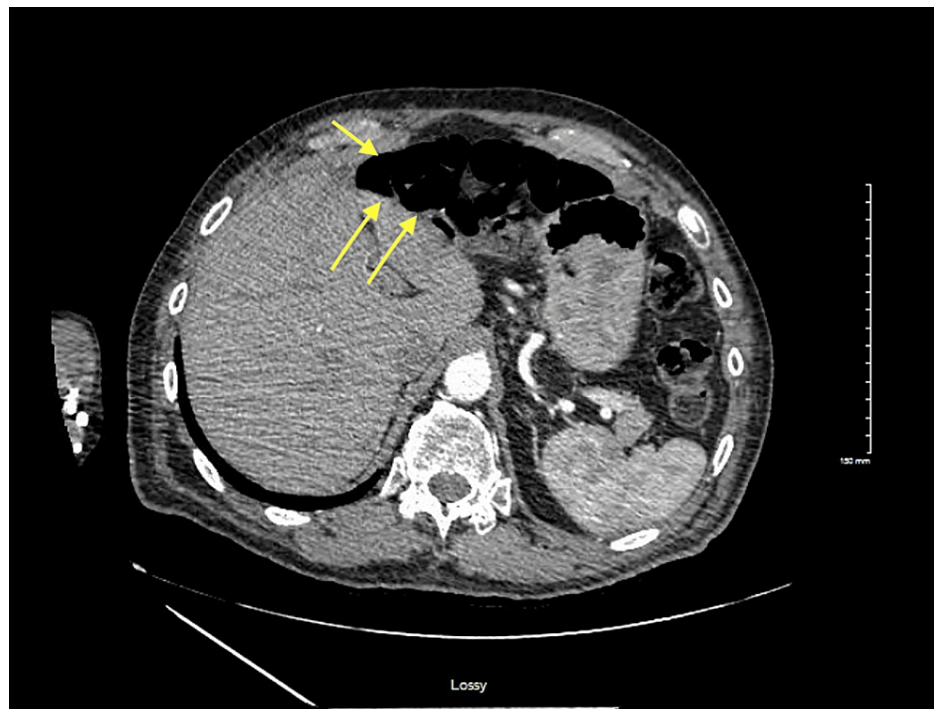


FIGURE 4: CT angiogram of the abdomen/pelvis

Abdominal CT angiogram reveals the bowel loops receding and not interposed between the liver and abdominal wall

CT: computed tomography

No acute intra-abdominal pathology was identified, and the patient improved with conservative management.

Discussion

On admission, our patient was found to be hypotensive and have lactic acidosis, concerning for an ischemic or infectious process and potential sepsis. Imaging was obtained and demonstrated subdiaphragmatic air. As previously mentioned, the etiologies of subdiaphragmatic lucency are broad and potentially emergent. This case underscores the importance of integrating radiologic findings with clinical status: employing surgical management could have increased morbidity for our patient, given his baseline frailty. The chart review demonstrated that the apparent air was a chronic process and not due to an acute pathology such as bowel ischemia and perforation, even with the elevated lactate. The patient was noted to have the anatomic variant of the Chilaiditi sign, a critical distinction that led to conservative treatment.

The prevalence of the Chilaiditi sign is highest in the elderly [10]. Considering the aforementioned drivers of the Chilaiditi sign, intestinal hypermobility and an enlarged subdiaphragmatic space, the reasons for this may be generalized frailty and colonic redundancy secondary to comorbid constipation, a condition that correlates positively with age [10,11]. We considered possible etiologies for our patient's Chilaiditi sign, and given his baseline muscle weakness, we proposed that incomplete diaphragmatic excursion could have led to subdiaphragmatic space enlargement. Meanwhile, chronic constipation may have contributed to colonic redundancy and displacement.

While our case had the benefit of prior images, an isolated finding of subdiaphragmatic air is often concerning. Therefore, it is important to consider imaging techniques that may increase suspicion of the Chilaiditi sign, as many emergency department patients will present without baseline examinations. Typical features include haustral markings that surround the subdiaphragmatic air and a lack of positional change of the subdiaphragmatic air when the patient moves [12]. This second characteristic is because free, extraluminal air is gravity-dependent, whereas the Chilaiditi syndrome air is contained by the interpositioned colon. Even if the patient is in the left lateral decubitus position, the Chilaiditi sign air will still be trapped in the subdiaphragmatic space. Such imaging pivot points would be of paramount importance in the presented case, as the patient's age, chronic weakness, frail state, and lactic acidosis would have made him a poor surgical candidate. Moreover, the high surgical risk is an issue likely common to many Chilaiditi syndrome patients, considering its elderly predilection [10].

The Chilaiditi syndrome has a diverse presentation, including respiratory distress and angina-like episodes [6]. The appropriate treatment therefore considers symptom severity, with critical manifestations such as those complicated by bowel obstruction prompting surgical intervention [13]. The chronicity and frequency of associated symptoms also have a role in dictating management. A recent case report proposed staging and treatment protocols for the Chilaiditi syndrome, with recommendations dependent on these factors [14]. Following their proposed algorithm for our case would have led to the same course of conservative treatment; however, such tools are useful in supporting management decisions and for settings in which the syndrome recurs. After inpatient stabilization, mild instances of the syndrome could also be aided by physical activity after discharge, as exercise could promote intestinal peristalsis and symptom resolution [15]. Meanwhile, for severe cases, additional workup could include Gastrografin enemas, CT with endoluminal contrast, and colonoscopy [14]. Reported successful surgical interventions for these instances include total or partial colonic resection combined with hepatoxy [6,13]. Some patients have even required diaphragmatic intervention [16,17].

Conclusions

In summary, the Chilaiditi sign, a rare radiologic finding caused by colonic interposition between the liver and diaphragm, can progress to the Chilaiditi syndrome when clinical symptoms arise. Despite its low incidence, computed tomography is more sensitive in detection. Its etiology involves congenital and acquired factors, often affecting elderly individuals. Differential diagnosis is crucial due to imaging manifestations resembling serious conditions. Our case highlights the importance of accurate diagnosis, as conservative management effectively resolved symptoms, avoiding surgery in a high-risk patient. Tailored treatment strategies, guided by symptom severity and recurrence, are essential, with recent literature proposing staging and treatment protocols. Understanding the diverse presentations and management options is critical for optimizing outcomes, especially in elderly patients with multiple comorbidities.

Additional Information

Author Contributions

All authors have reviewed the final version to be published and agreed to be accountable for all aspects of the work.

Concept and design: Ibrahim Kamel, Reid Ponder, Ibrahim Elkhawas, Yusuf Yalcin, Zeeshan Solangi

Acquisition, analysis, or interpretation of data: Ibrahim Kamel, Reid Ponder, Ibrahim Elkhawas, Yusuf Yalcin

Drafting of the manuscript: Ibrahim Kamel, Reid Ponder, Ibrahim Elkhawas, Yusuf Yalcin

Critical review of the manuscript for important intellectual content: Ibrahim Kamel, Reid Ponder, Ibrahim Elkhawas, Yusuf Yalcin, Zeeshan Solangi

Supervision: Zeeshan Solangi

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Farkas R, Moalem J, Hammond J: Chilaiditi's sign in a blunt trauma patient: a case report and review of the literature. *J Trauma*. 2008, 65:1540-2. [10.1097/01.ta.0000208194.49228.03](https://doi.org/10.1097/01.ta.0000208194.49228.03)
2. Prassopoulos PK, Raissaki MT, Gourtsoyiannis NC: Hepatodiaphragmatic interposition of the colon in the upright and supine position. *J Comput Assist Tomogr*. 1996, 20:151-3. [10.1097/00004728-199601000-00027](https://doi.org/10.1097/00004728-199601000-00027)
3. Murphy JM, Maibaum A, Alexander G, Dixon AK: Chilaiditi's syndrome and obesity. *Clin Anat*. 2000, 13:181-4. [10.1002/\(SICI\)1098-2355\(2000\)13:3<181::AID-CA4>3.0.CO;2-7](https://doi.org/10.1002/(SICI)1098-2355(2000)13:3<181::AID-CA4>3.0.CO;2-7)
4. Kang D, Pan AS, Lopez MA, Buicko JL, Lopez-Viego M: Acute abdominal pain secondary to chilaiditi syndrome. *Case Rep Surg*. 2013, 2013:756590. [10.1155/2013/756590](https://doi.org/10.1155/2013/756590)
5. Tariq HA, Pillay T: The air up there - Chilaiditi's syndrome: a case report and review of the literature. *Afr J Emerg Med*. 2020, 10:266-8. [10.1016/j.afjem.2020.04.001](https://doi.org/10.1016/j.afjem.2020.04.001)
6. Moaven O, Hodin RA: Chilaiditi syndrome: a rare entity with important differential diagnoses. *Gastroenterol Hepatol (N Y)*. 2012, 8:276-8.
7. Kamiyoshihara M, Ibe T, Takeyoshi I: Chilaiditi's sign mimicking a traumatic diaphragmatic hernia. *Ann*

- Thorac Surg. 2009, 87:959-61. [10.1016/j.athoracsur.2008.07.033](https://doi.org/10.1016/j.athoracsur.2008.07.033)
8. Vallee PA: Symptomatic Morgagni hernia misdiagnosed as Chilaiditi syndrome . West J Emerg Med. 2011, 12:121-3.
 9. Orangio GR, Fazio VW, Winkelman E, McGonagle BA: The Chilaiditi syndrome and associated volvulus of the transverse colon. An indication for surgical therapy. Dis Colon Rectum. 1986, 29:653-6. [10.1007/BF02560330](https://doi.org/10.1007/BF02560330)
 10. Yin AX, Park GH, Garnett GM, Balfour JF: Chilaiditi syndrome precipitated by colonoscopy: a case report and review of the literature. Hawaii J Med Public Health. 2012, 71:158-62.
 11. Schuster BG, Kosar L, Kamrul R: Constipation in older adults: stepwise approach to keep things moving . Can Fam Physician. 2015, 61:152-8.
 12. Joo YE: Chilaiditi's sign. Korean J Gastroenterol. 2012, 59:260-1. [10.4166/kjg.2012.59.3.260](https://doi.org/10.4166/kjg.2012.59.3.260)
 13. Kao CT, Dunkley M, Hodgson R: Surgical management of large bowel obstruction and significant hepatic displacement caused by Chilaiditi syndrome. BMJ Case Rep. 2023, 16:e255047. [10.1136/bcr-2023-255047](https://doi.org/10.1136/bcr-2023-255047)
 14. Adu Y, Nesiama EA, Siddiqui A, Prakash S, Obokhare I: Chilaiditi syndrome: a case report, literature review, and proposition of a novel management staging system. Cureus. 2023, 15:e46688. [10.7759/cureus.46688](https://doi.org/10.7759/cureus.46688)
 15. Xu Y, Wang Q, Meng G, et al.: A rare cause of sudden chest pain and dyspnea: a CARE-compliant case report of Chilaiditi syndrome. Medicine (Baltimore). 2020, 99:e20220. [10.1097/MD.0000000000002020](https://doi.org/10.1097/MD.0000000000002020)
 16. Richardson B, Hickham L, Harper S, Soliman B: Delayed right diaphragmatic hernia with Chilaiditi syndrome: a case report. Cureus. 2023, 15:e41420. [10.7759/cureus.41420](https://doi.org/10.7759/cureus.41420)
 17. Sidhu KK, Van Kessel CS, Cao C, Austin KK: The combination of Chilaiditi syndrome and Bochdalek hernia in an adult: successful management with a robot assisted approach. ANZ J Surg. 2023, 93:1035-7. [10.1111/ans.18068](https://doi.org/10.1111/ans.18068)