# Chilaiditi Sign: A Rare Radiological Finding

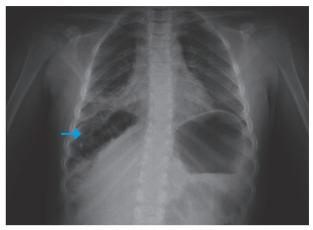
Pedro Mantas, Sara Machado Rocha, Margarida Marujo, Aldina Canteiro Lopes

Port J Pediatr 2021;52:70-1 DOI: https://doi.org/10.25754/pjp.2021.19571

A 7-year-old girl, with no relevant medical history and normal development, presented acutely with a six-day history of coughing and pyrexia. Physical examination was normal except for tympany on percussion over the right upper abdominal quadrant. A plain chest radiograph showed a middle lobe consolidation with possible rightsided pleural effusion and incidentally detected hepatodiaphragmatic interposition of the colon (Fig. 1), known as the Chilaiditi sign. The observed colonic haustration suggested that there was no free intraperitoneal air. Blood tests showed a rise in the inflammatory markers (leukocytes 10,900 cells/µL, with 57.1% neutrophils and 32.9% lymphocytes, C-reactive protein 9.14 mg/dL) and the blood culture was negative. Right pleural effusion was observed with thoracic ultrasound. The patient was admitted for observation and started on antibiotics for pneumonia. She remained stable throughout her admission, with no associated symptoms or worsening radiological changes and she was discharged home after three days.

First described in the medical literature in 1910 by Greek radiologist Demetrius Chilaiditi, the Chilaiditi sign refers to the rare incidental radiologic finding of the hepato-diaphragmatic interposition of the colon.<sup>1</sup> The worldwide incidence is estimated at 0.025-0.28%.<sup>2</sup> The etiology is multifactorial, but not fully understood.<sup>2</sup> Generally, patients are asymptomatic. When associated with symptoms, typically gastrointestinal or respiratory, it is designated as Chilaiditi syndrome.<sup>2,3</sup>

The clinical relevance of Chilaiditi sign and syndrome relate to a wide differential diagnosis that includes serious conditions such as pneumoperitoneum, diaphragmatic hernia, volvulus, and intussusception, requiring urgent surgical management.<sup>2-4</sup> Although no intervention is required for an asymptomatic patient and the initial management of Chilaiditi syndrome is conservative, patients should be kept under clinical and radiologic observation in order to prevent complications.<sup>5</sup> Despite predominance in adults,<sup>2-4</sup> a rather limited number of pediatric cases are reported in the literature, starting in the neonatal period<sup>2,3</sup> and, therefore, pediatricians should be aware of this condition.



**Figure 1.** Chest radiograph. Chilaiditi sign: right hepato-diaphragmatic interposition of the colonic haustration. Middle lobe consolidation. Absent visualization of right diaphragm suggesting pleural effusion.

Keywords: Chilaiditi Syndrome / diagnostic imaging; Child

#### WHAT THIS REPORT ADDS

- Chilaiditi sign is a rare radiologic finding of the hepato-diaphragmatic interposition of the colon.
- Generally, it is asymptomatic but can be associated with symptoms, typically gastrointestinal or respiratory.
- Differential diagnosis includes pneumoperitoneum, diaphragmatic hernia, volvulus, and intussusception.
- Initial management of Chilaiditi syndrome is conservative, with clinical and radiologic observation.

#### **Conflicts of Interest**

The authors declare that there were no conflicts of interest in conducting this work.

#### **Funding Sources**

There were no external funding sources for the realization of this paper.

#### Provenance and peer review

Not commissioned; externally peer reviewed

#### **Consent for publication**

Consent for publication was obtained.

### **Confidentiality of data**

The authors declare that they have followed the protocols of their work centre on the publication of patient data.

#### Awards and presentations

Poster presentation at the 9th Excellence in Pediatrics Conference, at Wien, Austria, December 2017

Pediatrics Department, Hospital de Santarém, Santarém, Portugal Corresponding Author Pedro Mantas

https://orcid.org/0000-0003-1741-3023

pedromantas@gmail.com

Avenida Bernardo Santareno, 37-37B, 2005-177 Santarém, Portugal

Received: 28/02/2020 | Accepted: 01/06/2020 | Published: 03/01/2021

© Author(s) (or their employer(s)) and Portuguese Journal of Pediatrics 2021. Re-use permitted under CC BY-NC. No commercial re-use.



## References

1. Chilaiditi D. Zur frage der hepatoptose und ptose im allgemeinen im Anschluss an drei falle von temporarer, partieller leberverlagerung. Fortcshr Geb Rontgenstr Nuklearmed Erganzongsband 1910;16:173-208.

2. Jiménez OC, Ávila MB, Montes EP, Córdoba JD, Camacho RV. Signo y síndrome de Chilaiditi: condiciones infrecuentes pero con importancia diagnóstica en pediatría. Casos clinicos. Rev Chil Pediatr 2017;88:635-9. doi: 10.4067/S0370-41062017000500010.

3. Evrengul H, Yüksel S, Orpak S, Özhan B, Agladioglu K.

Chilaiditi syndrome. J Pediatr 2016;173:260. doi: 10.1016/j. jpeds.2016.02.060.

4. Moaven O, Hodin RA. Chilaiditi syndrome: A rare entity with important differential diagnoses. Gastroenterol Hepatol 2012;8:276-8.

5. Barroso Jornet JM, Balaguer A, Escribano J, Pagone F, Domenech J, del Castillo D. Chilaiditi syndrome associated with transverse colon volvulus: First report in a paediatric patient and review of the literature. Eur J Pediatr Surg 2003;13:425-8. doi: 10.1055/s-2003-44737.

