

Journal of Clinical Surgery and Research

Saad Slaiki *

Open Access

Case Report

The Hidden Breast: A Compelling Case of Supernumerary Breast

Saad Slaiki 1*, Jihad Jamor 2

¹ Visceral surgery department, CHU Hassan II, Morocco, Fes.

*Corresponding Author: Saad Slaiki, Visceral surgery department, CHU Hassan II, Morocco, Fes.

Received Date: 10 March 2023 | Accepted Date: 24 March 2023 | Published Date: 25 April 2023

Citation: Slaiki S., Jamor J., (2023), The Hidden Breast: A Compelling Case of Supernumerary Breast, *Journal of Clinical Surgery and Research*, 4(2); **DOI:**10.31579/2768-2757/072

Copyright: © 2023, Saad Slaiki. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract:

Supernumerary breast is a rare condition in which one or more additional breasts develop along the mammary line. Although it is often asymptomatic, it can cause discomfort or lactation problems in some women. This article with a review of literature provides a comprehensive overview of supernumerary breast, including its definition, prevalence, symptoms, diagnosis, treatment, consequences, and prevention.

keywords: polymastia; supernumerary; breast

Introduction

Axillary supernumerary breast tissue, also known as polymastia, is a rare condition characterized by the presence of extra breast tissue in the axilla [1]. While diagnosing supernumerary breasts is relatively straightforward in the presence of a nipple and lactating discharge, it becomes challenging in their absence or when there is a significant amount of adipose tissue [2]. In this article, we will provide a brief overview of axillary supernumerary breast tissue, its clinical presentation, and its diagnosis and management, based on the available literature. We will also discuss the importance of recognizing this condition and the need for further research to better understand its prevalence and clinical significance.

Case presentation

A Young 28-year-old woman with no notable medical history, who sought medical attention due to the presence of a left axillary mass that had been gradually increasing in size for a year, particularly around the time of her menstrual cycle. Physical examination revealed a well-defined, soft, painless, and mobile mass measuring 9 cm in diameter that was adherent to the skin but not to the deep plane. There were no visible skin changes around the mass, and the initial diagnosis was an axillary lipoma (figure 1).



Figure 1: left supernumerary breast

² Gynecology department, CHU Hassan II, Morocco, Fes.

The rest of the physical exam was unremarkable. An ultrasound was performed and revealed ectopic glandular tissue surrounded by fatty tissue in the left axillary fossa without any detectable nodular or cystic lesion (figure 2).



Figure 2: ultrasound picture suspecting Supernumerary Breast

Surgical excision was performed (figure 3), and the anatomopathological study confirmed the presence of a supernumerary breast with no evidence of tumor cells. The patient had an uneventful postoperative recovery.



Figure 3: Resected Specimens

Discussion

Supernumerary breast, also known as polymastia or accessory breast, is a rare congenital condition characterized by the development of one or more additional breasts along the milk line [3]. This condition occurs in approximately 1-5% of the population, with the highest prevalence in females [1]. It's generally considered to be benign, but it can sometimes cause discomfort or embarrassment for the patient. Furthermore, axillary supernumerary breast tissue can be confused with other benign or malignant conditions, such as lipomas or cysts, making diagnosis and management challenging [2].

The etiology of supernumerary breast is not fully understood, but it is believed to be caused by the persistence of embryonic mammary ridges, which give rise to breast tissue during fetal development [4].

The diagnosis of supernumerary breast is made through physical examination and imaging studies, such as mammography, ultrasound, and magnetic resonance imaging (MRI) [5]. In some cases, a biopsy may be

necessary to confirm the diagnosis [6]. Once a diagnosis has been made, the management of axillary supernumerary breast tissue depends on the symptoms and clinical presentation of the patient. If the condition is causing discomfort or cosmetic concerns, surgical excision may be considered [5].

Although axillary supernumerary breast tissue is a rare condition, it is important for clinicians to be aware of its existence and the potential for misdiagnosis. Misdiagnosis can lead to unnecessary procedures and delayed treatment. Therefore, accurate diagnosis and appropriate management of axillary supernumerary breast tissue can improve patient outcomes and avoid unnecessary interventions.

Further research is needed to better understand the prevalence and clinical significance of axillary supernumerary breast tissue. Studies have shown that the condition can be difficult to diagnose and may be misdiagnosed as other benign or malignant conditions [7]. More research is needed to improve diagnostic accuracy and develop evidence-based management guidelines.

Conclusion

Axillary supernumerary breast tissue is a rare condition that can present diagnostic challenges. However, with appropriate diagnosis and management, patients can be effectively treated and their symptoms alleviated. Healthcare providers should be aware of this condition and consider it in the differential diagnosis of patients presenting with axillary masses.

References

- Grossl NA. (2000). Supernumerary breast tissue: historical perspectives and clinical features. South Med J. 93(1):29-32.
- 2. Margi M, Azhary O, Oulahyane R, Cherkaoui A, Abdelhak M, et al. (2010). Sein surnuméraire axillaire: à propos d'un cas. *Archives de Pédiatrie*. 17(8):1162-1164.
- 3. Weinshel LR, Demakopoulos N. (1943). Supernumerary breasts: With special reference to the pseudomamma type. *The American Journal of Surgery*. 60(1):76-80.

- Garba A, Adamou H, Amadou Magagi I, Hassane DM, Mamane D. (2017). Polymastie axillaire bilatérale: une observation familiale. Bilateral axillary polymastia: a familial case. Annales africanes de médecine, 10:2572-2575.
- Singal R, Mehta SK, Bala J, Zaman M, Mittal A, et al. (2016).
 A Study of Evaluation and Management of Rare Congenital Breast Diseases. J Clin Diagn Res. 10(10): PC18-24.
- Hiremath B, Subramaniam N, Chandrashekhar N. (2015). Giant accessory breast: a rare occurrence reported, with a review of the literature. BMJ Case Rep.
- Almatrafi MI, Almalki MA, Althagafi JA, AlSindi TS, Masarit RM, et al. Benign Breast Disease in Makkah, Saudi Arabia: A Retrospective Analytical Cross-Sectional Study. *Cureus*. 14(11): e31174.



This work is licensed under Creative Commons Attribution 4.0 License

To Submit Your Article Click Here:

Submit Manuscript

DOI: 10.31579/2768-2757/072

Ready to submit your research? Choose Auctores and benefit from:

- > fast, convenient online submission
- > rigorous peer review by experienced research in your field
- > rapid publication on acceptance
- authors retain copyrights
- unique DOI for all articles
- immediate, unrestricted online access

At Auctores, research is always in progress.

Learn more https://www.auctoresonline.org/journals/journal-of-clinical-surgery-and-research