Case Report

A Case of Granulomatosis with Polyangiitis (GPA) Where a Multicystic Nasal Septal Abscess Aided in the Diagnosis

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1. Introduction

Granulomatosis with polyangiitis (GPA), formerly known as Wegener’s granulomatosis, is a systemic refractory vasculitis that presents with necrotising granulomatous inflammation in the upper respiratory tract and lungs, necrotising glomerulonephritis in the kidney, and necrotising vasculitis of medium and small arteries throughout the body. GPA is a systemic refractory vasculitis. Although the prognosis of this disease is poor, early diagnosis and therapeutic intervention have led to the remission of the disease in an increasing number of cases. Physical examination is important in the diagnosis of granulomatosis polyangiitis. Upper respiratory tract symptoms are often reported as saddle nose and perforation of the nasal septum, and only one case of the nasal septal abscess has been reported as an upper respiratory tract symptom.

Herein, we describe a case of GPA with the nasal septal abscess that resulted in a visit to our department. Since upper respiratory tract symptoms account for 70%–100% of all cases of GPA, a patient will likely visit an otorhinolaryngologist for the first time. The nasal cavity and paranasal sinuses are the most common sites involved in the head and neck region (85%–100%), but ear symptoms may develop initially in approximately 35% of cases, as discussed by Greco [1].

Otorhinolaryngologists need to be aware of GPA, which often develops due to upper respiratory tract symptoms. However, the actual diagnosis is not easy because lesions confined to the upper respiratory tract alone often do not present with symptoms and findings typical of GPA.

There have been many reports of nasal involvement due to GPA, such as thickening of the mucosa of the nasal septum [2], perforation of the nasal septum [3], saddle nose
due to nasal bone destruction [4], and chronic dacryocystitis and nasolacrimal duct stenosis [5]. However, there have been few reports of nasal septal abscesses, such as in our case. Moreover, there are few reports of nasal septal abscesses, and only two cases, including an autopsy, have been reported.

2. Case Report/Case Presentation

A 69-year-old male patient presented to the outpatient clinic of the Department of Otorhinolaryngology with a chief complaint of nasal obstruction persisting for 4 months. He had a history of diabetes mellitus and no history of rheumatoid arthritis.

Physical examination revealed swelling of the anterior nasal septum and dorsum of the nose (Figures 1(a)-1(b)). Blood tests showed that the white blood cell count was elevated to $15.8 \times 10^3$, and C-reactive protein had increased to 11.76. Meanwhile, the creatinine level was in the normal range, and there were no proteinuria, haematuria, or rheumatological problems. Chest computed tomography (CT) also showed no abnormality. Contrast-enhanced CT showed multiple abscesses in the nasal septum (Figures 2(a)-2(b)).

Drainage surgery was performed in parallel with histopathological examination. Intraoperative findings showed white necrosis between the nasal septum cartilage and mucosa, as well as white necrosis and pus accumulation in the periosteum in the pear-shaped opening to the nasal bone. The mucosa of the nasal septum was incised and drained, and a biopsy of the mucosa in and surrounding the abscess was performed endoscopically (Video 1).
Bacterial abscess formation was suspected; therefore, antimicrobial agent therapy was initiated immediately after the surgery, albeit with no improvement. Subsequently, blood tests revealed a prolonged inflammatory response and nasal septal abscess culture tests were negative during hospitalisation. Further blood tests showed PR3-ANCA positivity. In addition, epithelial cell granuloma and vasculitis were discovered in the specimens collected during emergency surgery. From this information, we reached a consensus regarding our diagnosis. After confirming the prognosis, we immediately started the patient on steroids at 40 mg per day due to an early systemic form of granuloma, resulting in an improvement in the swelling of the anterior nasal septum and dorsum of the nose and the indented oedema of the dorsum of both hands, and the fever receded.

Consequently, intravenous cyclophosphamide 500 mg per 2 weeks was also added as remission induction therapy.

In addition, blood tests revealed an improvement in the inflammatory response. One month after the surgery, the swelling of the external nose subsided, followed by a saddle nose, a typical physical finding of granulomatosis with GPA (Figures 3(a)-3(b)). The patient remains in good condition and is being carefully monitored in the outpatient clinic.

3. Discussion/Conclusions

The probability of obtaining characteristic histological findings on biopsy for GPA is only 40%–60% [2, 6]. Therefore, the following technique for biopsy tissue collection is required. Areas of strong ulceration or necrosis must be avoided, and a large sample of an adjacent undisturbed area must be taken (specifically, more than 5 mm in diameter) [7]. Samples from multiple sites must be taken and the procedure must be repeated if there is no specific finding [6].

When GPA is considered or suspected, an active biopsy must be performed [6, 8, 9].

In our case, a nasal septal abscess was the main symptom, which is not an indicator for the diagnosis of GPA; therefore, it was difficult to consider it in the differential diagnosis.

However, we affirmed that this case was atypical because the contrast-enhanced CT scan showed multifocal abscess formation in the nasal septum. Due to this abnormality, we performed a biopsy for diagnostic purposes.

Of the five cases of nasal septal abscess confirmed by contrast-enhanced CT in previous studies, three were bacterial [10–13], one was fungal [13], and all four were monoculartype. Meanwhile, the other cases caused by GPA showed multifocal findings, as in our case [14].

In conclusion, when an investigation reveals a multifocal nasal septal abscess, the possibility of GPA should be considered as an atypical case of infectious abscess formation.

Data Availability

The data are not publicly available due to privacy and ethical restrictions.

Ethical Approval

All procedures were performed in accordance with the ethical standards of the institutional and/or National Research Committee and adhered to the tenets of the 1964 Declaration of Helsinki and its later amendments or comparable ethical standards.

Conflicts of Interest

The authors declare that there are no conflicts of interest.

Authors’ Contributions

MS: collected the data and wrote the manuscript. KO, TE, and NO: edited the manuscript.
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Supplementary Materials

Video 1. Intraoperative endoscopic findings. The nasal septal mucosa is biopsied from different sites on both sides to prevent nasal septal perforation (biopsied from low horizontal on the left and high horizontal on the right septum membrane). A concha bullosa is also biopsied as a relatively normal area. (Supplementary Materials)

References

[1] A. Greco, C. Marinelli, M. Fusconi et al., “Clinic manifestations in granulomatosis with polyangiitis,” *International Journal of Immunopathology & Pharmacology*, vol. 29, no. 2, pp. 151–159, 2016.

[2] J. Ohori, “Granulomatosis with polyangiitis,” *Journal of Japan Society of Immunology & Allergology in Otolaryngology*, vol. 37, no. 4, pp. 265–268, 2019.

[3] Coordes, S. M. Loose, V. M. Hofmann et al., “Saddle nose deformity and septal perforation in granulomatosis with polyangiitis,” *Clinical Otolaryngology*, vol. 43, no. 1, pp. p291–p299, 2017.

[4] F. Bello and F. Fagni, “Nasal deformity in granulomatosis with polyangiitis,” *New England Journal of Medicine*, vol. 382, no. 10, p. 951, 2020.

[5] B. Jiang, Y. Y. Zhao, and S. H. Wei, “Granulomatosis with polyangiitis: the relationship between ocular and nasal disease,” *Ocular Immunology and Inflammation*, vol. 21, no. 2, pp. 115–118, 2013.

[6] A. Masiak, Z. Zdrojewski, R. Pęksa et al., “The usefulness of histopathological examinations of non-renal biopsies in the diagnosis of granulomatosis with polyangiitis,” *Reumatologia/Rheumatology*, vol. 5, no. 5, pp. 230–236, 2017.

[7] E. A. Del Buono and A. Flint, “Diagnostic usefulness of nasal biopsy in Wegener’s granulomatosis,” *Human Pathology*, vol. 22, no. 2, pp. 107–110, 1991.

[8] O. Matsubara, N. Yoshimura, Y. Doi, A. Tamura, and E. J. Mark, “Nasal biopsy in the early diagnosis of Wegener’s (pathergic) granulomatosis. Significance of palisading granuloma and leukocytoclastic vasculitis,” *Virchows Archiv*, vol. 428, no. 1, pp. 13–19, 1996.

[9] S. Srinivasan, S. Begum, S. Kathirvelu, and A. Vaithy, “Limited granulomatosis with polyangiitis presenting as an isolated lung lesion,” *Indian Journal of Pathology & Microbiology*, vol. 63, no. 4, pp. 611–614, 2020.

[10] M. Berlucchi, M. Tomasoni, R. Bosio, and V. Rampinelli, “Spontaneous abscess of the posterior nasal septum: an unusual cause of Nasal Obstruction in children,” *Annals of Otology, Rhinology & Laryngology*, vol. 130, no. 8, pp. 966–969, 2021.

[11] Y. Hyo, H. Fukushima, T. Harada, and H. Hara, “Nasal septal abscess caused by anaerobic bacteria of oral flora,” *Auris Nasus Larynx*, vol. 46, no. 1, pp. 147–150, 2019.

[12] Y.-C. Huang, P. L. Hung, and H. C. Lin, “Nasal septal abscess in an immunocompetent child,” *Pediatrics & Neonatology*, vol. 53, no. 3, pp. 213–215, 2012.

[13] J. M. Debnam, A. M. Gillenwater, and L. E. Ginsberg, “Nasal septal abscess in patients with immunosuppression,” *American Journal of Neuroradiology*, vol. 28, no. 10, pp. 1878-1879, 2007.

[14] I. Spanuchart, N. Zungsontiporn, P. O-charoen, B. Changcharoen, and D. T. Bolger, “Granulomatosis with polyangiitis: a case of nasal mass, necrotic lung, and normal kidneys,” *Hawai’i Journal of Medicine & Public Health: A Journal of Asia Pacific Medicine & Public Health*, vol. 74, no. 9 Suppl 2, pp. 27–29, 2015.