

A Midline Nasal Mass in a Term Neonate



A term infant with pneumothorax was admitted to the neonatal unit for respiratory support after birth. Examination revealed a skin-colored mass extending from the nasal septum, occluding the right nasal aperture (**Figure 1**). The pneumothorax resolved, but the respiratory distress persisted with desaturations, particularly during feeding. A nasopharyngeal airway was inserted.

A postnatal computed tomography scan showed a 3-cm peripherally enhancing mass completely occluding the right nasal passage extending to the crista galli (**Figure 2**). Assessment of the bony anatomy revealed no intracranial extension. On day 14, the lesion was debulked endoscopically. Histopathological analysis confirmed a nasal glioma.

Congenital nasal anomalies are rare, affecting only 1 in 20 000-40 000 newborns.¹ Resulting from disordered development of the primitive neural plate, types include dermoids, encephaloceles, and gliomas.² Nasal gliomas are thought to be encephaloceles, with 20% having a vestigial stalk connecting to the intracranial space.³ Gliomas are typically located proximal to the nasal root. Histologically, they resemble encephaloceles but are often quite vascular and easily mistaken for hemangiomas.² Neuroimaging can help determine the presence or absence of intracranial connections.⁴ Biopsy should be avoided until intracranial attachment has been excluded.

Left untreated, nasal gliomas can result in bony atrophy and intracranial communications can act as inoculation sites with a risk of meningitis and encephalitis. Treatment is complete surgical excision early in life to avoid complications and achieve optimal cosmetic outcomes. ■

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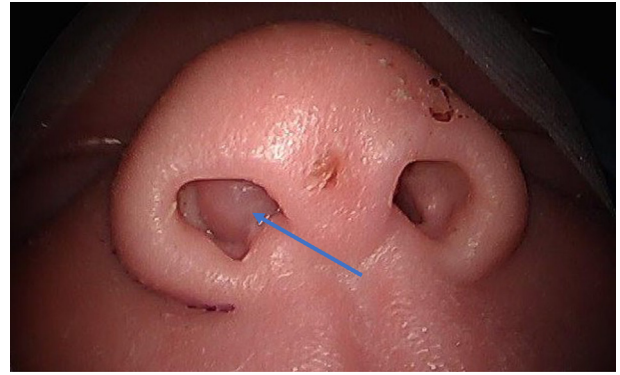


Figure 1. Nasal mass obstructing the right naris.

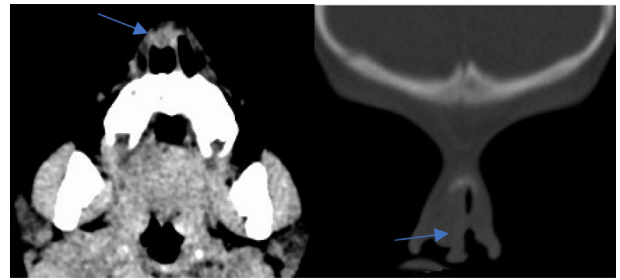


Figure 2. Noncontrast computed tomography scan showing a soft tissue lesion in the anterior right nasal passage.

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