Self-induced ethmoidectomy from rhinotillexomania.

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Self-induced Ethmoidectomy from Rhinotillexomania


Summary: A 53-year-old woman with a long history of compulsive nose picking (rhinotillexomania) presented with a large, self-inflicted nasal septal perforation and right-sided penetration of the ethmoidal sinus, or “ethmoidectomy.”

Index terms: Nose, injuries; Paranasal sinuses

Chronic self-mutilation resulting in the loss of body parts is characteristically seen in schizophrenic patients. Such patients can have delusions of parasitic infestation of body parts, may believe the body part to be encumbered by foreign bodies, or may view the body part as no longer a part of themselves (1). Such behavior, however, may also be manifested by persons who are severely obsessive-compulsive or malingerers (1).

Case Report

A 53-year-old right-handed woman related a history of compulsive nose picking (rhinotillexomania) of the right nasal cavity since age 10. She could not control her compulsion, which involved removing recurrent intranasal crusts. This condition persisted while in the care of a psychiatrist. When referred to an otolaryngologist, a large right-sided self-induced ethmoidectomy cavity and a large nasal septal perforation from the trauma were noted. There was no history of nasal surgery, nor was an incision present. Computed tomography (CT) revealed the dilated ethmoidal cavity to be in proximity to both the orbit and the cribiform plate (Fig 1). Therapy was instituted in an effort to disrupt the cycle of digital trauma, mucus production, and crusting. This included behavior modification and supportive rhinologic care with nasal spray, crust suction, and medication. Early follow-up showed improvement.

Discussion

Serious upper airway destruction due to compulsive nose picking is not addressed in standard textbooks of otolaryngology and pediatrics, although nose picking resulting in epistaxis and/or septal perforation is cited (2–5). Conversely, the psychiatric literature has recognized that rhinotillexomania is a common, benign habit in children and adults that may rarely become a serious affliction advancing to significant self-injury (6). Such injuries are known to include nasal septal perforation and epistaxis, which may be resistant to control because of repetitive trauma (7–9). Anemia from the chronic blood loss may ensue, and subjects may attempt to conceal their affliction from their physician.

Interestingly, there are few prior reports of massive injury. Akhtar and Hastings (1) described a 36-year-old man, a compulsive nose picker, who had life-threatening epistaxis. Self-destructive behavior was so apparent that the left ala of the nose was absent, exposing the nasal passage. His injury was not the result of a true psychosis, but of a severe “passive-aggressive character disorder.” Gigliotti and Waring (10) reported a 61-year-old woman with extensive self-mutilation affecting portions of the nasal septum, turbinates, hard palate, and other internal nasal structures such that a nasal prosthesis and complete upper denture had to be constructed. Neither of these case reports included radiologic studies.

The differential diagnosis in our patient would include granulomatous disease, such as Wegener granulomatosis, or cocaine abuse, although advanced unilateral involvement would be unusual.

In conclusion, we offer this uncommon case, with CT documentation, to demonstrate the extent of nasal and paranasal sinus destruction that may occur from self-inflicted digital nasal trauma in persons who are obsessive-compulsive or schizophrenic. The possibility of self-
inflicted injuries should be considered whenever such findings are encountered.

References
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