

Abstract

Background: Sparse data are available in literature about pattern of nasal affection in Sjogren syndrome (SS).

Objective: To study the ultrastructural changes in the nasal mucosa in SS patients with nasal dryness.

Methods: Light microscopy and transmission electron microscopy (TEM) of anterior end of inferior turbinate in 14 patients with SS experiencing nasal dryness and five healthy controls who underwent turbinate reduction surgery were included in this study. Nasal symptoms were assessed according to visual analogue score. Patients were subjected to nasal endoscopy and computerized tomography of paranasal sinuses when indicated.

Results: Light microscopy showed mild to severe lymphocytic infiltration of nasal submucosa with dilated ducts, sparse seromucinous acini and mild fibrosis. The overlying epithelium showed variable squamocolumnar hyperplasia or atrophy with prominent goblet cell depletion. The basal lamina zone appeared apparently thickened and irregular with hyalinosis. TEM revealed disorganized surface epithelium. The basal lamina was frequently very thin atrophic and breached. Dense collagen bundles occupied the submucosa. Collagen bundles frequently extended through breached basal lamina (BM) to the surface epithelium in a process resembling cirrhosis. Blood vessels showed vasculitis. Sparse seromucinous glands showed minimal mucin and apoptotic myoepithelial and glandular cells.

Conclusion: This study is the first to describe ultra structural changes of nasal mucosa in SS, especially nasal cirrhosis. Ultrastructural changes were generally indicative of an underlying autoimmune process and may add to better understanding of pathophysiology of SS. Lastly nasal affection in Sjogren syndrome is underestimated.