



NTP Nonneoplastic Lesion Atlas

Tongue, Epithelium – Hyperplasia



Figure Legend: Figure 1 Tongue, Epithelium - Hyperplasia in a female F344/N rat from a chronic study. A focus of epithelial hyperplasia is present on the tongue (arrow). **Figure 2** Tongue, Epithelium - Hyperplasia in a female F344/N rat from a chronic study (higher magnification of Figure 1). There are papillary epithelial projections in the focus of epithelial hyperplasia.

Comment: Squamous cell hyperplasia in the oral cavity is seen most commonly on the tongue, palate, and lateral wall of the pharynx. Squamous cell hyperplasia is characterized by increased cell numbers, which usually results in increased thickness of the squamous epithelium. Squamous hyperplasia may be diffuse or plaque-like or may form blunt papillary projections. Focal hyperplasia can have multiple finger-like projections, each with its own lamina propria (Figure 1 and Figure 2). Hyperkeratosis is frequently seen with squamous cell hyperplasia, but usually keratin pearl formation is not present. Size, the well-differentiated appearance of cells, and the absence of a prominent stromal component are important in distinguishing hyperplasia from papilloma. Papillomas frequently will have a single stalklike attachment to the tongue, which may not be in the plane of section. Proliferative lesions involving squamous cell carcinoma. There are, however, examples of studies in which the papilloma stage was apparently bypassed, and progression was directly from focal hyperplasia to squamous cell carcinoma. Mechanical damage can result in ulceration or erosion of the hyperplastic epithelium and inflammation.



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Recommendation: Squamous hyperplasia should be diagnosed and graded based on the size and

number of the areas affected and the thickness of the epithelium. Associated hyperkeratosis,

ulceration, or inflammation should not be diagnosed separately unless warranted by severity.

References:

Brown HR, Hardisty JF. 1990. Oral cavity, esophagus and stomach. In: Pathology of the Fischer Rat (Boorman GA, Montgomery CA, MacKenzie WF, eds). Academic Press, San Diego, CA, 9-30. Abstract: <u>http://www.ncbi.nlm.nih.gov/nlmcatalog/9002563</u>

Leininger JR, Jokinen MP, Dangler CA, Whiteley LO. 1999. Oral cavity, esophagus, and stomach. In: Pathology of the Mouse (Maronpot RR, eds). Cache River Press, St Louis, MO, 29-48. Abstract: <u>http://www.cacheriverpress.com/books/pathmouse.htm</u>

National Toxicology Program. 2010. NTP TR-544. Toxicology and Carcinogenesis Studies of Dibromoacetonitrile (CAS No. 3252-43-5) in F344/N Rats and B6C3F1 Mice (Drinking Water Studies). NTP, Research Triangle Park, NC. Abstract: http://ntp.niehs.nih.gov/go/32617

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