

## Oral Histopathology

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### Series 42 (11 cases)

Case	Features
Fibrolipoma	<ul style="list-style-type: none"><li>Nodule composed largely of adipose tissue with some fibrous connective tissue intervening</li></ul>
Peripheral giant cell granuloma	<ul style="list-style-type: none"><li>Gingival nodule with multinucleated giant cells</li></ul>
Dentigerous cyst	<ul style="list-style-type: none"><li>Impacted tooth with simple cyst lining; the connective tissue in the cyst wall has a fibrous to myxoid appearance; in those cases where lining cannot be identified, the terminology <i>fibromyxomatous connective tissue consistent with the wall of a dentigerous cyst</i> may be used</li></ul>
Odontogenic keratocyst	<ul style="list-style-type: none"><li>Palisading (alignment) of basal nuclei, 5-8 cell layers, and corrugated parakeratin</li></ul>
Adenomatoid odontogenic tumor	<ul style="list-style-type: none"><li>Swirls and gland-like structures of epithelioid to spindle shaped odontogenic epithelium with some ghost cells, calcifications and dentin-like material</li><li>Often the clinical presentation is the "lesion of 2s or 2/3's" – 2/3 anterior, 2/3 maxilla, 2/3 female and 2/3 in the first 2 decades of life; presenting as a radiolucency with or without radiopacities</li></ul>
Schwannoma	<ul style="list-style-type: none"><li>A benign neural (nerve-related) lesion characterized by <i>Antoni A</i> or <i>Antoni B</i> tissue (the Antoni A tissue consists of palisaded <i>Verocay bodies</i>)</li><li>This case is a combination of Antoni A and Antoni B tissue and consists of streams of bland nerve tissue with elongated but plump nuclei with some nuclear inclusions but no marked pleomorphism or mitoses</li></ul>
Neurofibroma	<ul style="list-style-type: none"><li>Juxtapose this to the schwannoma, this lesion consists of spindled nerve cells with comma-shaped or wavy nuclei in a fibrous background stroma</li><li>The presence of mast cells (not identified here) also suggests the diagnosis of <i>neurofibroma</i></li></ul>
Verruciform xanthoma	<ul style="list-style-type: none"><li>Clinically these are papillary lesions which overlap with squamous papillomas and in some cases papillary dysplasias</li><li>Histology shows V-shaped/chevron/papillary/verruccous epithelium with an orange-red color to the keratin which 'plugs' the epithelium and the characteristic foam-like <i>xanthoma cells</i> present in the papillary connective tissue</li></ul>
Seborrheic keratosis	<ul style="list-style-type: none"><li>Clinically this is the waxy/raisin-like 'stuck on' lesion, often on the face, often in individuals in their 4<sup>th</sup>-5<sup>th</sup> (or older) decades of life</li><li>There is epidermal thickening, formation of keratin pearl-like cysts, a flat to pushing basal layer of cells, and there may be pigmentation</li></ul>
Blue nevus	<ul style="list-style-type: none"><li>Clinically these are usually blue/gray to brown macules present on skin or mucosa</li><li>The histology shows spindle-shaped highly pigmented cells in the dermis/submucosa which often run parallel to the overlying epithelium</li></ul>
Melanoma	<ul style="list-style-type: none"><li>The topic of melanoma is complex and often requires the input of a skin pathologist</li><li>The basic features that lead to the diagnosis in this case are variably sized nests (theques) with cellular and nuclear atypia and pleomorphism; pigment may be present (as in this case) or absent; there may also be <i>Pagetoid</i> (vertical) spread of abnormal melanocytes through the epithelium</li></ul>