

## Oral Histopathology

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### Series 24 (18 cases)

Case	Features
Benign mixed tumor	<ul style="list-style-type: none"><li>Well-defined partially encapsulated tumor (palate) consisting of myoepithelial cells, some hyalinized and myxoid stroma, duct formation and some areas of keratinization and calcification (no two tumors ever look truly alike!)</li><li>Adjacent minor salivary glands noted</li></ul>
Fibroma, giant cell type	<ul style="list-style-type: none"><li>A fibroma, often seen on tongue and gingiva, often with a papillary appearance (DDX includes papilloma) with dense collagen and large stellate or triangular shaped fibroblasts, some with multiple nuclei (<i>giant cell fibroma</i> is not to be confused with <i>giant cell granuloma</i>)</li></ul>
Ameloblastoma	<ul style="list-style-type: none"><li>This case has both cystic and solid/plexiform morphology, palisading of basal nuclei, reverse nuclear polarity and stellate reticulum formation (treatment was as for a solid, not cystic, ameloblastoma)</li></ul>
Melanoma	<ul style="list-style-type: none"><li>This is a poorly differentiated malignancy with marked cellular and nuclear pleomorphism, numerous abnormal mitoses (some are star-shaped); the presence of brown melanin pigment and large purple to dark 'cherry red' nucleoli aided in diagnosis, as did immunohistochemistry (S-100, Mart-1 and HMB-45, stains for melanocytes) and a clinical history of previous melanoma</li></ul>
Peripheral odontogenic fibroma, WHO type	<ul style="list-style-type: none"><li>Gingival mass consisting of fibrous connective tissue and numerous epithelial odontogenic rests</li></ul>
Peripheral ossifying fibroma, ulcerated	<ul style="list-style-type: none"><li>Ulcerated gingival nodule with dystrophic calcifications and granulation tissue underlying the ulcer</li></ul>
Squamous papilloma	<ul style="list-style-type: none"><li>Papillary gingival nodule; the apparent separation of the individual papillary growths is an artifact of processing and cutting across the papillary 'fronds'</li></ul>
Lobular capillary hemangioma, ulcerated	<ul style="list-style-type: none"><li>Surface is ulcerated and covered in an eosinophilic fibrin; the underlying tissue mass is composed of lobules of vascular tissue (small capillaries)</li><li>Academically, one can debate the diagnosis of <i>pyogenic granuloma</i> as neither 'pyogenic' (i.e. induced by fever) or 'granuloma' (i.e. granulomatous inferring a foreign body reaction or true granulomas as seen in sarcoidosis or tuberculosis) or if a better diagnostic terminology is, as in this case <i>lobular capillary hemangioma</i> or <i>granulation tissue type hemangioma</i> [terminology can be confusing, therefore the different diagnostic terms are introduced here in context to aid the participant/reader]</li></ul>
Reaction to polarizable material (lip augmentation)	<ul style="list-style-type: none"><li>Polarizable foreign material and associated multinucleated foreign body type giant cells</li><li>Reader is referred to the earlier series on foreign body materials as an exercise to determine the exact nature of the material</li><li>A library of foreign materials is also available at the AAOMP web site, <a href="http://www.aaomp.org/">http://www.aaomp.org/</a> - scroll to the ATLAS for the pull down menu foreign materials library</li></ul>
Pulse granulomas	<ul style="list-style-type: none"><li>The terms <i>pulse granuloma</i>, <i>connective tissue hyaline bodies</i> and <i>giant cell hyaline angiopathy</i> are sometimes used</li></ul>

	<p>interchangeably (but probably should not be)</p> <ul style="list-style-type: none"> <li>• In this case, a polarizable fragment (vegetable material or 'pulse') is identified so the term 'pulse granuloma' is most appropriate</li> <li>• Involvement of blood vessels may favor the term 'hyaline angiopathy' and a simple hyaline mass with giant cells (as in some areas of this case) may favor the term 'connective tissue hyaline body'</li> </ul>
Leukoedema	<ul style="list-style-type: none"> <li>• Rarely biopsied, since diascopy or stretching of the bluish-white 'leukoplakia' of buccal mucosa (usually in darker skinned individuals) leads to blanching or disappearance of the lesion and is satisfactory for diagnosis</li> <li>• Likely represents a defect in production of certain keratins</li> <li>• Evident histologically, as in this case, as 'ballooning' and acanthosis (thickening) of the epithelial cells in the spinous layer</li> <li>• Frictional keratosis may mimic this histologically (look for surface maceration and bacterial debris)</li> <li>• Oral hairy leukoplakia is almost identical histologically; a history of immune suppression (solid organ transplant, HIV/AIDS) will aid in diagnosis and immunohistochemistry or genetic hybridization for Epstein Barr virus or its component genes/proteins may also be employed</li> </ul>
Squamous cell carcinoma, well to moderately differentiated	<ul style="list-style-type: none"> <li>• Most of this carcinoma is keratinizing (well differentiated) but other areas take on a more spindle cell characteristic and less keratin production (moderately differentiated)</li> <li>• Other features are still present (cellular and nuclear pleomorphism, prominent nucleoli, keratin pearl formation, carcinoma arising/dropping from surface epithelium which is highly dysplastic and dyskeratotic)</li> </ul>
Mucocele, extravasation type	<ul style="list-style-type: none"> <li>• Mucus surrounded by compressed granulation tissue and partially by a ruptured duct epithelium</li> <li>• Minor salivary gland lobules without significant pathologic changes</li> </ul>
Mucoepidermoid carcinoma	<ul style="list-style-type: none"> <li>• Classic features: numerous mucocytes (mucus cells), epidermoid and intermediate cells; predominantly solid (rather than cystic) but still histologically 'bland'</li> </ul>
Benign fibro-osseous lesion, c/w focal COD	<ul style="list-style-type: none"> <li>• Abnormally shaped bone and osteocementum, round spherical cementicles and a spindle cell to vascular background [the radiograph showed a mixed radiodensity adjacent to but not attached to a molar root]</li> </ul>
Ameloblastoma	<ul style="list-style-type: none"> <li>• Basal palisading, reverse polarity, stellate reticulum</li> <li>• Some areas show <i>granular cell</i> and <i>acanthomatous (squamous)</i> changes in the stellate reticulum</li> <li>• Other areas show connective tissue fibrosis and 'compression' of the tumor nests and strands in the so-called <i>desmoplastic</i> variant [this is a unique variant which appears radiologically as a mixed radiodensity and may mimic a BFOL]</li> </ul>
Ameloblastoma	<ul style="list-style-type: none"> <li>• Predominantly cystic architecture but otherwise shows the characteristic features (basal palisading, reverse polarity, stellate reticulum which here lines the cyst lumen [in these lesions these are known as the <i>Vickers-Gorlin criteria</i> and delineate these cystic ameloblastomas from other cysts])</li> </ul>
Odontogenic keratocyst	<ul style="list-style-type: none"> <li>• Basal palisading (without reverse polarity, compare to the previous two cases), 4-8 cell layers, corrugated parakeratin</li> </ul>