

## Oral Histopathology

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### Series 28 (13 cases)

Case	Features
Orthokeratinizing odontogenic cyst	<ul style="list-style-type: none"> <li>Only mild hint at basal palisading but marked orthokeratosis (nuclei lacking in keratin layer, prominent granular layer)</li> </ul>
Odontogenic keratocyst	<ul style="list-style-type: none"> <li>Basal palisading, 8+ layers of cells, corrugated parakeratin</li> </ul>
Odontogenic keratocyst, cortical plate	<ul style="list-style-type: none"> <li>Same case as previous, showing significant thinning of the cortical plate</li> </ul>
Odontogenic keratocyst	<ul style="list-style-type: none"> <li>Basal palisading, 8+ layers of cells in some areas (or some transverse sectioning creating artifactual 'thickening' of the cyst lining), parakeratin</li> </ul>
Salivary duct cyst	<ul style="list-style-type: none"> <li>An epithelial lined cyst adjacent to minor (mucus) glands on the palate; the thickening and apparent separation of the cyst lining from the connective tissue may represent artifact and orientation more than true epithelial proliferation</li> </ul>
Salivary duct cyst with oncocytic metaplasia	<ul style="list-style-type: none"> <li>Single to double layer of oncocytic (pink) cells line this cyst</li> </ul>
Squamous cell carcinoma with necrosis	<ul style="list-style-type: none"> <li>Well-defined keratinizing squamous cell carcinoma with nuclear pleomorphism, prominent nucleoli, keratin pearl formation; some areas show necrosis (the 'architecture' is still present but the cells and nuclei are lost in a sea of amorphous keratin)</li> <li>Necrosis is considered a factor for poor prognostic outcome</li> </ul>
Squamous cell carcinoma, perineural and perivascular invasion	<ul style="list-style-type: none"> <li>Similar histology to previous case, but the carcinoma is seen abutting a large/muscular artery and surrounding/abutting a nerve in the higher magnification views</li> <li>Lymphovascular and perineural invasion are considered factors for poor prognostic outcomes</li> </ul>
Adenoid cystic carcinoma	<ul style="list-style-type: none"> <li>This carcinoma shows the variability in morphology that may be seen; some areas show cribriform ("Swiss cheese") with basement membrane like material in the pseudocystic space, other areas are more tubular; nuclei are generally small, darker and irregular to wedge-shaped</li> </ul>
Peripheral ossifying, ulcerated	<ul style="list-style-type: none"> <li>Gingival nodule with bone and osteoid, ulcer and fibrin</li> </ul>
Neurofibroma	<ul style="list-style-type: none"> <li>Ill-defined proliferation of spindle cells admixed with neural to fibrous stroma, nuclei are comma or wavy in shape</li> <li>Numerous mast cells ("purple fried eggs") are a common finding in neurofibromas and lipid (fat) tumors</li> </ul>
Granulomatous inflammation (Crohn's disease)	<ul style="list-style-type: none"> <li>Inflammation and multiple granulomas (accumulations of epithelioid histiocytes and multinucleated giant cells); there is a tissue fold in one area (an artifact of processing)</li> <li>Special stains (PAS, GMS, acid fast bacillus) were negative; patient had significant GI disease</li> </ul>
Sequestrum, c/w mature BFOL (trephined implant)	<ul style="list-style-type: none"> <li>A mature osteocementum with reversal lines was trephined with a failing implant; the fibrillary to amorphous lavender material is bacterial debris</li> </ul>