

MICROSCOPIC DESCRIPTION: Nasal mucosa (per contributor – full credit for oral mucosa or lip, but it's not tongue – not enough skeletal muscle) **(1 pt)**: Expanding the nasal submucosa, there are numerous well-formed pyogranulomas **(2 pt)**. Pyogranulomas are centered on surrounded by radiating, club-like brightly eosinophilic material **(1 pt)** (Splendore-Hoepli material) **(2 pt)** and are composed of numerous viable and degenerate neutrophils **(1 pt)** admixed with cellular debris and more peripherally, numerous epithelioid macrophages **(1 pt)**, few lymphocytes and plasma cells **(1 pt)**, enmeshed in tight lamellations of fibrous connective tissue. **(1 pt)** There is diffuse fibrosis **(1 pt)** of the submucosa between pyogranulomas which extends into the underlying skeletal muscle. Entrapped skeletal muscle fibers demonstrate increased eosinophilia, shrinkage (atrophy) **(1 pt)** and mild hypertrophy of satellite nuclei. Low numbers of neutrophils, macrophages, lymphocytes and plasma cells are spread thinly throughout the fibrotic submucosa **(1 pt)**, and salivary gland tissue contains few lymphocytes and plasma cells within the interstitium. **(1 pt)**

MORPHOLOGIC DIAGNOSIS: Nasal mucosa (or lip) **(1 pt)**: Rhinitis , pyogranulomatous **(1 pt)**, chronic, multifocal to coalescing, moderate, with Splendore-Hoepli material **(1 pt)**.

Cause: *Actinobacillus lignieresii* **(2 pt)**

O/C: **(1 pt)**

**(Note: If you described bacilli in the colonies of Splendore-Hoepli material, they aren't visible on the HE (or even a Gram stain.)**

Kidney: Diffusely, there are changes at all levels of the tubule. Approximately 20% of tubules at all levels within the cortex and less commonly, the medulla (**1pt**) contain sheaves or fan-like arrangements of birefringent translucent crystals (**2pt**) (oxalates) (**2pt**), which occasionally rupture the basement membrane (**1pt**). Ruptured tubules are surrounded by loosely arranged collagen and low numbers of lymphocytes and plasma cells. (**1pt**) In many crystal-laden tubules, lining epithelium demonstrates a range of morphologic changes: swelling and vacuolation (degeneration) (**1pt**), shrunken with pyknosis and sloughed into the lumen (necrotic) (**1pt**), and some tubules are lined by attenuated epithelium and contain luminal protein (**1pt**). Bowman's space is filled with granular reflux and there is hypertrophy of parietal epithelium. (**1pt**) There are three empty subcapsular cysts, the largest being 1.7mm in diameter.

Cerebrum: Perivascular areas within the meninges and extending down along Virchow-Robin spaces are multifocally expanded by clear space (edema) (**1pt**). Birefringent crystals are present within both vessel lumina and in perivascular areas. (**1pt**)

MORPHOLOGIC DIAGNOSIS: 1. Kidney, tubules: Degeneration and necrosis (**1pt**), diffuse, marked , with marked numerous intratubular oxalate crystals (**1pt**).  
2. Cerebrum, vessels: Rare intramural oxalate crystals. (**1pt**)

CAUSE: Ethylene glycol toxicosis, (primary oxalosis OK). (**3 pt.**)

O/C: (**1 pt**)

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Conf. 4, Case 3.

Tissue from an ox.

**MICROSCOPIC DESCRIPTION:** Haired skin: Diffusely within the superficial dermis there is a perivascular, periadnexal, and perifollicular (**1pt.**) inflammatory infiltrate composed of large numbers of eosinophils (**1pt.**) and macrophages (**1pt.**) with fewer neutrophils, lymphocytes and plasma cells. (**1pt.**) Multifocally, hair follicles are dilated and contain at their base, several cross and tangential sections of follicle of adult (**1pt.**) filarial (**1pt.**) nematodes that are 100 um in diameter (**1pt.**) with a 5 um thick smooth cuticle, polymyarian-coelomyarian musculature (**1pt.**), a pseudocoelom, lateral alae, a thick walled intestine lined by uninucleate cuboidal cells, and reproductive organs including paired uteri containing microfilariae (**1pt.**) and eosinophilic discs (female) or a testis (male). (**1pt.**) At least one follicle is ruptured, with adult nematodes free in the dermis where they (and liberated keratin debris) incite a profound inflammation reaction of neutrophils and macrophages. (**1pt.**) Multifocally, hair follicles are ectatic, lined by attenuated epithelium, and filled with lamellations of keratin and occasionally degenerate neutrophils and necrotic debris (luminal folliculitis) (**1pt.**); the follicular epithelium is also occasionally infiltrated by neutrophils (mural folliculitis). Rarely follicles are surrounded by dense fibrous connective tissue and are hyperplastic and tortuous. (**1pt.**) There is mild orthokeratotic hyperkeratosis (**1pt.**) and diffuse epidermal hyperplasia (**1pt.**) characterized by rete ridge formation, acanthosis, prominent intercellular bridging (spongiosis), and intracellular clear space (intracellular edema). There are multifocal intracorneal pustules (**1pt.**) that contain degenerate neutrophils, cellular debris, acantholytic keratinocytes, and proteinaceous fluid. Multifocally there are serocellular crusts (**1pt.**) containing degenerate neutrophils, serum, cellular debris, hemorrhage, and entrapped plant material and bacterial colonies.

**MORPHOLOGIC DIAGNOSIS:** Haired skin: Dermatitis, perifollicular, periadnexal and perivascular, (**1pt.**) eosinophilic (**1pt.**) and histiocytic, diffuse, moderate, with folliculitis, furunculosis, dermal microfilariae, and few intrafollicular adult filarial nematodes (**1pt.**)

**CAUSE:** *Stephanofilaria stilesi* (**3pt.**)

**O/C:** (**1pt.**)

Case 4. Tissue from a cynomolgus macaque.

MICROSCOPIC DESCRIPTION: Pancreas. Diffusely throughout the pancreas, islets are increased both in number (**2pt.**) and in size (**2pt.**), ranging up to .5mm in diameter. Within affected islets, islet cells are hypertrophied with marked anisokaryosis (**1pt.**), prominent nucleoli and occasionally form acini. Many, if not all islets are infiltrated by low to moderate numbers of lymphocytes (**2pt.**) which separate islet cells and occasionally form large aggregates (**1pt.**), effacing parts of the islets. There are occasionally single necrotic cells surrounded by lymphocytes in infiltrated islets. (**1pt.**) The vast majority of islets are further expanded by a variable accumulation of a waxy, hyaline material (**1pt.**) (amyloid) (**2pt.**) that separates, compresses and often replaces islet cells, infiltrating lymphocytes, and minimally extends into the surrounding parenchyma. Islet cells in amyloid-effaced islets often have vacuolated cytoplasm (degeneration) (**1pt.**) or are shrunken and pyknotic (necrosis) and/or mineralized. (**1pt.**). Acinar cells are generally unremarkable but occasionally have clear lipid vacuoles in their cytoplasm.

MORPHOLOGIC DIAGNOSIS: 1. Pancreas, islets: Hyperplasia (**1pt.**), diffuse, moderate with islet cell hypertrophy(**1pt.**).

2. Pancreas, islets of Langerhans: Amyloidosis, diffuse, severe. (**1pt.**)

3. Pancreas, islets of Langerhans Insulitis, lymphocytic, multifocal, moderate. (**1pt.**)

NAME A LIKELY CLINICOPATHOLOGIC ABNORMALITY: Hyperglycemia (1pt.)

O/C: (**1pt.**)