

WSC 2021-2022

Conference 7, Case 1.

Tissue from a dog.

MICROSCOPIC DESCRIPTION: Cerebrum: Effacing (**1pt.**) 80% of the neuroparenchyma is a poorly circumscribed, unencapsulated, poorly demarcated, multilobular, infiltrative neoplasm (**1pt.**) The neoplasm is composed of spindle cells (**1pt.**) loosely arranged into long interlacing streams and bundles (**1pt.**) on a fine vascular stroma. (**1pt.**) Neoplastic cells have indistinct cell borders and a moderate amount of vacuolated eosinophilic cytoplasm. (**1pt.**) Nuclei are oval to elongate with coarsely stippled chromatin and 2-3 small blue nucleoli. (**1pt.**) There is moderate anisocytosis and anisokaryosis, (**1pt.**) and mitotic figures average 10 per 2.37mm^2 field. (**1pt.**) There are multiple large, serpiginous and occasionally coalescing areas (**1pt.**) in which there is architectural retention with loss of differential staining (necrosis), with mild multifocal infiltrates of neutrophils, cellular debris, and hemorrhage (1pt), fibrin and edema. (**1pt.**) Vessels in affected areas are also necrotic and often contain occlusive or non-occlusive thrombi and extramural hemorrhage. (**1pt.**) Surrounding these necrotic areas are perpendicularly oriented pseudopalisades of neoplastic cells (**1pt.**) There are numerous multilayered microvascular proliferations (**1pt.**) present in tortuous or glomeruloid arrangements. There is marked gliosis and mild spongiosis of the neuroparenchyma surrounding the neoplasm and widely scattered spheroids. (**1pt.**)

MORPHOLOGIC DIAGNOSIS: Cerebrum: Astrocytoma, high grade. (Astrocytoma grade 4 (glioblastoma multiforme old, but okay.) (**3pt.**)

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

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Conference 7, Case 2.

Tissue from a cat.

MICROSCOPIC DESCRIPTION: Rostral cerebrum (**1pt.**): Arising from the meninges (**1pt.**), and compressing (**1pt.**) the adjacent neocortex, there is an unencapsulated, moderately cellular, well-demarcated, nodular neoplasm (**1pt.**). Neoplastic cells are arranged in short interlacing streams (**1pt.**), bundles, and whorls (**1pt.**) on a moderate fibrovascular matrix (**1pt.**). Neoplastic cells are spindled (**1pt.**) with indistinct cell borders and a moderate amount of a finely fibrillar cytoplasm (**1pt.**). Nuclei are oval with finely clumped chromatin and 1-2 prominent basophilic nuclei. (**1pt.**) Mitotic figures are rare. (**1pt.**) Neoplastic cells have distinct whorling around vessels or granular to spherical aggregates of densely mineralized eosinophilic matrix. (**1pt.**) There are numerous areas of aggregated foamy macrophages, (**1pt.**) large acicular cholesterol clefts (**1pt.**), and less commonly, crystalline mineral. At the advancing front of the neoplasm, the adjacent compressed neuropil is loosely arranged, with mild edema and mild astrocytic gliosis. (**1pt.**)

MORPHOLOGIC DIAGNOSIS: Cerebrum: Meningioma (**4pt.**) (No classification schema for cats yet).

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

MICROSCOPIC DESCRIPTION: Brainstem, colliculi, cerebellum (**2pt.**): Multifocally, numerous brainstem neurons (**2pt.**), as well as neurons within all layers of the cerebellum (sparing granule cells) (**1pt.**) contain a single or multiple 10-40 um diameter (**3pt**) round structure with a blue-gray, often lamellated core (**2pt**) surrounded by a 2-4 um wide lightly amphophilic, finely granular peripheral zone (**1pt.**) and further outlined by a 1um thin zone of eosinophilic material (peripheralized axoplasm) (Lafora bodies) (**2pt.**). Purkinje cells occasionally contain multiple Lafora bodies. (**1pt.**) Lafora bodies are seen both within axons (**1pt.**) and less commonly in the nuclear perikarya. (**1pt.**) There is a mild diffuse increase in microglia in regions in which the preponderance of neurons contain inclusions (**1pt.**)kk and multifocal areas of mildly spongiotic white matter in the brainstem.

MORPHOLOGIC DIAGNOSIS: Brainstem, cerebellum: Intraneuronal Lafora (polyglucosan) bodies (**1pt.**) numerous, with mild scattered microgliosis (**1pt.**) and spongiosis.

NAME THE CONDITION: Lafora body disease. (**2pt.**)

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

Case 4. Tissue from a dog.

MICROSCOPIC DESCRIPTION: Multiple sections, cerebrum (at level of caudate nucleus), cerebellum, and brainstem. Within the cerebral gray and white matter (**1pt.**), but more prominently in the white matter, the neuropil is diffusely infiltrated by an unencapsulated, moderately cellular, poorly demarcated neoplasm (**1pt.**) composed of neoplastic astrocytes (**2pt.**). Neoplastic cells are arranged singly or in small clusters (**1pt.**) infiltrating the pre-existing neuroparenchyma. Neoplastic cells primarily appear as “naked nuclei” (**1pt.**) or with scant amounts of poorly defined finely granular eosinophilic cytoplasm (**1pt.**). Nuclei are pleomorphic, with most being elongate but others ovoid, irregular, or round, with finely stippled hyperchromatic chromatin and a 1-3 visible nucleoli. (**1pt.**) . (**1pt.**) Anisocytosis and anisokaryosis are regionally variable (**1pt.**), with areas in which there are 2-4 per standardized 400x field. (**1pt.**) In areas of dense infiltration by neoplastic cells within areas of white matter, neoplastic cells are separated by amphophilic granular debris (axonal debris?). (**1pt.**) Neoplastic cells often surround neurons, vessels, or submeningeal areas in small clusters (secondary structures) (**1pt.**) Similar changes are multifocally present in the brainstem; the cerebellum appears to be spared. (**1pt.**) Astrocytes with large nuclei, marginated chromatin, prominent nucleoli, and abundant eosinophilic cytoplasm (gemistocytes) are present in areas of neoplastic invasion.

MORPHOLOGIC DIAGNOSIS: Cerebrum, brainstem: Astrocytoma, infiltrative (diffuse OK), high grade (previously known as gliomatosis cerebri). (**4pt.**)

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