

Clinical classification of neuronal ceroid-lipofuscinosis

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Citation Report

#	ARTICLE	IF	CITATIONS
1	CEROID“LIPOFUSCINOSIS (BATTEN'S DISEASE): PATHOGENESIS AND SEQUENTIAL NEUROPATHOLOGICAL CHANGES IN THE OVINE MODEL. <i>Neuropathology and Applied Neurobiology</i> , 1989, 15, 371-383.	3.2	68
2	Infantile Osteopetrosis Complicating Neuronal Ceroid Lipofuscinosis. <i>Pathology Research and Practice</i> , 1990, 186, 697-706.	2.3	16
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8	Jansky-Bielschowsky variant disease: CT, MRI, and SPECT findings. <i>Pediatric Neurology</i> , 1992, 8, 121-126.	2.1	57
9	Abnormal lysosomal cathepsin activities in leukocytes and cultured skin fibroblasts in late infantile, but not in juvenile neuronal ceroid-lipofuscinosis (Batten disease). <i>Clinica Chimica Acta</i> , 1992, 208, 111-117.	1.1	10
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14	Linkage analysis in juvenile neuronal ceroid lipofuscinosis. <i>American Journal of Medical Genetics Part A</i> , 1992, 42, 542-545.	2.4	10
15	Seizures, depression and dementia in teenagers with Batten disease. <i>Journal of Inherited Metabolic Disease</i> , 1993, 16, 252-255.	3.6	21
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