

EA1 - kirjallisuusviitteet

Bressman S: Genetics of movement disorders: recent advances. Kokousyhteenvedossa: Movement disorders, s. 440-67 – 440-102. American Acadedy of Neurology, 48th Annual Meeting, March 23 – 30, San Fransisco, 1996

Brandt T and Strupp M: Episodic ataxia type 1 and 2 (familial periodic ataxia/vertigo). Audiol Neurotol 2 (6): 373 – 383, 1997

Kaakkola S ja Rinne R: Ataksiat ja niiden erotusdiagnostiikka. Duodecim 113: 1773 – 1782, 1997

Meisler MH et al., Ion channel mutations in mouse models of inherited neurological disease. Ann Med 29: 569 – 574, 1997

Somer H ja Auranen M: Lihastautien geenivirheet. – katsaus. Duodecim 113: 1795 – 1802, 1997

Barchi RL: Ion channel mutations affecting muscle and brain. Curr Opin Neurol 11: 461 – 468, 1998

Gordon N: Episodic ataxia and channelopathies. Brain&Development 20: 9 – 13, 1998

Subramony SH and Nance M: Diagnosis and management of the inherited ataxias. The Neurologist 4: 327 – 338, 1998

Davies NP and Hanna MG: Neurological channelopathies: diagnosis and therapy in the new millennium. Ann Med 31: 406 – 420, 1999

Brunt ER: Episodic ataxia type 1. Kirjassa: Handbook of Ataxia Disorders, 487 - 515. Ed. T Klockgether, Marcel Dekker Inc., New York-Basel, 2000

Eunson LH et al., Clinical, genetic, and expression studies of mutations in the potassium channel gene KCNA1 reveal new phenotypic variability. Ann Neurol 48(4): 647 – 656, 2000

Baloh RW: Periodic and progressive ataxias, sivut 229 - 238. Kirjassa: Channelopathies of the Nervous System. Toim. Rose MR ja Griggs RC. Butterworth and Heinemann, 2001

Goadsby PJ and Ferrari MD: Migraine: a multifactorial, neurovascular episodic channelopathy?, sivut 274 – 292. Kirjassa: Channelopathies of the Nervous System. Toim. Rose MR ja Griggs RC. Butterworth and Heinemann, 2001

Jen J (Ed): Ataxia and Calcium Channels. What a Headache! Arch Neurol 58: 179 – 180, 2001