Peripheral neuropathy in a family with Sandhoff disease and SH3TC2 deficiency

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Dear Sirs,

Sandhoff disease is a lysosomal storage disease caused by a deficiency of beta-hexosaminidase. Affected individuals present with a wide spectrum of clinical manifestations, ranging from psychomotor impairment and death in the infantile form to motor neuron disease and autonomic dysfunction in the adult form [1, 2]. Here, we present a family with predominantly sensory neuropathy that was found to have both Sandhoff disease and deficiency of SH3TC2.

A 46-year-old man had weakness, atrophy, and paresthesia with loss of sensation in his hands and feet for 30 years (Fig. 1a). He also had severe diarrhea several times per week, decreased sweating over his feet, and episodes of lightheadedness. On examination, he was found to have marked upper and lower extremity weakness and muscle atrophy distally, with the legs more severely affected (Fig. 1b). Although hyperreflexia was found in the upper extremities, reflexes in the lower extremities were reduced. Electromyography showed evidence of chronic neurogenic changes, and nerve conduction studies showed normal motor conduction velocities (patient 40–67 m/s, sibling 39–58 m/s). However, sensory nerve conduction studies were non-responsive in the proband and either non-responsive or reduced in amplitude and velocity (41 m/s) in the sibling. Autonomic testing (QSWEAT, Heart Rate Response to Deep Breathing, Valsalva, Tilt Table) and quantitative sensory testing (Vibratory Detection Threshold and Cooling Detection Threshold) showed evidence of autonomic dysfunction consistent with a severe small unmyelinated fiber and large myelinated fiber sensory neuropathy (Fig. 1d) [3]. Laboratory testing for acquired causes of neuropathy was negative. An evaluation of the proband’s 57-year-old sister showed a similar pattern of sensory, motor, and autonomic involvement, with symptom onset in her 20s. Both individuals did not show evidence of macular abnormalities; however, mild splenomegaly was found in the sister (span 14.2 cm).

After negative genetic testing (Athena Diagnostics Inc.) of ten candidate neuropathy genes (PMP22, Cx32, MPZ, PMP22, EGR2, NFL, PRX, GDAP1, LITAF, MFN2), whole exome sequencing (NISC/NHGRI/NIH) of the proband identified a common 1250C>T mutation (P417L) in HEXB and a previously reported [4] 2860C>T mutation (R954X) in SH3TC2, which was confirmed by Sanger sequencing (GeneDx). Further testing through multiplex ligation-dependent probe amplification (MLPA, GeneDx) showed that...
the proband is also heterozygous for a 16-kb deletion that includes exons 1–5 of the HEXB gene. The three variants were also confirmed in the affected sister, and although two of the unaffected sisters were found to have the SH3TC2 mutation, neither have both HEXB mutations. MR spectroscopy of the superior cerebellar vermis detected elevated creatine in the proband (Fig. 1c). Serum testing in both affected siblings showed low total hexosaminidase activity of 1.4 and 1.0 nmol/min/mL (normal 10.4–23.8) and an increased fraction of HEXA activity at 91 and 100 % (normal 56–80 %). A sural nerve biopsy from the proband showed evidence of a severe mixed axonal and demyelinating neuropathy (Fig. 1e–i) and a reduction in SH3TC2 immunoreactivity (Fig. 1j).

The adult form of beta-hexosaminidase deficiency is characterized by upper and lower motor neuron dysfunction, dysautonomia, and ataxia. The amino acid change in HEXB at position 417 is a common gene mutation in adult Sandhoff disease patients [5]. The early and severe presentation of sensory findings in both of these siblings is not a typical presentation of the disease [6–8]. The nonsense mutation observed in this family is predicted to reduce the accumulation of GM2 ganglioside in HEXB-deficient endosomes and lysosomes may be preferentially damaging to tissues that also have a deficiency of SH3TC2.

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Conflicts of interest The authors declare that they have no conflict of interest.

Ethical standard All human and animal studies have been approved by the appropriate ethics committee and have therefore been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments.

Informed consent All persons gave their informed consent prior to their inclusion in the study.

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