Deleterious mutation in $GPR88$ is associated with chorea, speech delay, and learning disabilities

ABSTRACT

Objective: To identify the underlying molecular basis of a familial developmental disorder characterized by chorea, marked speech delay, and learning difficulties in 4 sisters from a consanguineous family.

Methods: Whole-exome analysis of DNA of the 2 older patients followed by Sanger sequencing of the mutated exon in all family members.

Results: A homozygous deleterious mutation, p.C291X, was identified in the $GPR88$ gene in both exome analyses. The mutation segregated with the disease in the family and was absent from a large cohort of controls.

Conclusions: Homozygous deleterious mutation in $GPR88$ in humans is associated with marked speech delay, learning disabilities, and chorea, which manifest at 8–9 years of age. The finding is consistent with the reported abundant expression of GPR88 in the striatum and the hyperkinetic activity and learning impairment observed in GPR88 knockout mice. Although further functional characterization is needed, the finding underscores the importance of GPR88 in movement control and learning. *Neurrol Genet 2016;2:e64; doi: 10.1212/NXG.0000000000000064*

GLOSSARY

| ExAC | Exome Aggregation Consortium; GPext | globus pallidus pars externa; GPint | globus pallidus pars interna; MAF | minor allele frequency; MSN | medium spiny neuron. |

Chorea, defined as involuntary movements of limbs, trunk, neck, or face that rapidly flit from region to region in an irregular, flowing, nonstereotyped pattern, may be underlined by genetic, pharmacologic, metabolic, or anatomic causes. A practical approach to genetic childhood chorea is consideration of the mode of inheritance.1 X-linked chorea is seen in Lesch-Nyhan and Rett syndromes, and maternally inherited chorea is part of Leigh and MELAS syndromes. Autosomal dominant transmitted forms of chorea are usually part of a more complicated disorder, such as GLUT1 deficiency or PRRT2 mutation. A relatively “pure” chorea transmitted in an autosomal dominant mode is “benign hereditary chorea,” attributed to mutations in the $TITF1$ gene.2 Autosomal recessive transmitted forms of chorea are also part of a complex neurologic presentation, as in inborn errors of organic and amino acid metabolism, lysosomal storage diseases, mitochondrial respiratory chain defects, and brain iron accumulation (reviewed in reference 1).

We now report the results of the molecular investigation of patients from a consanguineous family who presented in childhood with choreiform movements, speech delay, and learning disabilities.

METHODS

Patients. Patient II-1 was the first daughter of first-cousin Palestinian parents (figure, A). Global developmental delay was noted in early infancy. She started walking at 24 months and uttered her first words at 3 years. Abnormal involuntary movements were first noted at the age of 9. These were initially mild and intermittent and confined to the face, predominantly around the mouth, as twitching of the lips. Within several months, the movements became more persistent and spread to the fingers and thighs.
had only subtle finger movements and intermittent upward shrug, occasional forward head movements were noted at 8 years. She uttered at 3 years of age. Repetitive rising of the eyebrows and grimacing, side-to-side movement of the pelvis and trunk, voluntary movements of the thighs and the head were also observed but at a lesser frequency. Motor impersistence of the tongue was noted. The rest of the neurologic examination was unremarkable. At 15 years, patient II-1 could read fluently but had marked dyscalculia. Her formal IQ was estimated at 40.

Patient II-4 had essentially a similar presentation and course. Psychomotor delay was evident in infancy; she walked independently at 3.5 years, presented in infancy with developmental delay, and at 8 years, she was unable to read or calculate. Neurologic examination was otherwise normal.

Patient II-8 was an 18-month-old girl at the time of writing. The pregnancy and delivery were uneventful, and her gross motor development was normal; she could sit at 8 months and walked independently at 18 months. However, at that age, there were no words at all. Abnormal movements were not seen.

Routine laboratory investigations including a battery of tests for inborn errors of metabolism were normal in patient II-1. This patient also underwent brain MRI at 14 years of age, which was normal.

The other children in the family (individuals II-2, II-3, II-6, and II-7) aged 3.5–13.5 years had normal motor development and no involuntary movements. All had age-appropriate fluent speech and average learning performance.

**Exome analysis.** Exome analysis was performed as previously described in the DNA samples of patients II-1 and II-4.

**RESULTS** The exome analyses of the DNA of patients II-1 and II-4 yielded 47.9 and 61.0 million reads, respectively (mean coverage X72.1 and X91.9). After alignment to the reference genome (Hg19) and variant calling, we removed variants that were called less than X8; were off-target, synonymous, or heterozygous; those with minor allele frequency (MAF) >1% in the Exome Aggregation Consortium (ExAC), Cambridge, MA (http://exac.broadinstitute.org); and those with MAF >2% in the Hadassah in-house database. Four homozygous variants were shared by the 2 sisters, but using Sanger sequencing we found that only 1 of the variants, Chr1:101005395C→T, NM_022049:c.C873A, p.C291X in the GPR88 gene, was carried by the parents and was found to be homozygous in patients II-5 and II-8. None of the other siblings, whose development was age appropriate at 3.5–13.5 years, was homozygous for the variant (figure, A and B). The variant was not carried by any of the ~60,000 individuals whose exome analyses were deposited in ExAC (accessed June 2015).

**DISCUSSION** The 4 female patients, ages 1.5–15 years, presented in infancy with developmental delay, marked speech retardation, and learning disability. The older 3, when reaching 8–9 years of age, developed movement disorder, which consisted of choreiform movements involving mainly the face and hands with involuntary movements of the shoulders, pelvis, and thighs. Molecular investigation revealed a homozygous nonsense mutation, p.C291X, in the GPR88 gene.

**GPR88** encodes a 384–amino acid protein, an orphan G protein–coupled receptor that is abundantly expressed in the striatal medium spiny neurons and at 8 years, she was unable to read or calculate. Neurologic examination was otherwise normal.
(MSNs). The MSNs play a crucial role in movement control. The motor cortex receives input from other cortical areas, the cerebellum, and the basal ganglia, always through the thalamus. The thalamus is under inhibitory activity of the globus pallidus pars interna (GPint); this inhibition can be released when the striatal GABAergic MSN transmits an inhibitory signal to the GPint. The inhibition is further relieved by stimulation of the MSN dopamine (D1) receptors. This signaling is the "direct pathway," which promotes movement. The opposing "indirect pathway," aims at preventing movement by using another striatal subpopulation of MSNs, which transmit an inhibitory signal to the globus pallidus pars externa (GPext). The inhibition is reinforced by stimulation of the MSN dopamine (D2) receptors. The GPext transmits inhibitory signal to the subthalamic nucleus, which in turn makes excitatory connections to cells in the GPint, maintaining the GPint inhibitory effect on the thalamus. The abundant expression of GPR88 in both D1 and D2 receptor–expressing MSNs suggests that GPR88 has an instrumental role in movement control. In agreement, our patients who were homozygous for a deleterious GPR88 mutation had a hyperkinetic movement disorder and the GPR88 knockout mice had increased locomotion, nonhabitual hyperactivity in novel environment (which is a primary landmark of striatal dysfunction), increased circling, and prolonged duration of burying episodes, suggesting stereotypic behavior. In the mutant animals, the MSNs had higher average firing, which was attributed to decreased tonic GABAergic inhibition resulting from low level of β3 protein, a GABAA receptor subunit, and to an increased glutamatergic excitatory synaptic transmission proposed to result from enhanced phosphorylation of the AMPA-type glutamate receptor subunit GluR1.6 Speech delay and poor school performance in our patients recapitulated the marked impairment in motor skill learning of the mutant mice despite normal coordination and strength. This deficit was confined to striatal learning and memory and did not affect hippocampal learning behavior.8 We propose that GPR88 deficiency in humans manifests as developmental delay with pronounced speech acquisition impairment, learning disabilities, and hyperkinetic movement disorder at 8–9 years of age. Although further functional studies are needed, our clinical data corroborate those observed in the GPR88 knockout mice and underscore the role of GPR88 in movement control and learning.

AUTHOR CONTRIBUTIONS
Dr. Fadi Alkufri and Dr. Bassam Abu-Libdeh designed and conceptualized the study and drafted and revised the manuscript. Dr. Avraham Shaag designed the study, analyzed and interpreted the data, and drafted the manuscript. Dr. Orly Elpeleg designed the study, analyzed and interpreted the data, and drafted and revised the manuscript.

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