Mutation-in-Brief

Novel AVPR2 variant in a male infant with nephrogenic diabetes insipidus who showed delayed head control

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Introduction

Congenital nephrogenic diabetes insipidus (NDI) is a rare disease caused by genetic mutations in AVPR2 or AQP2 (1). AVPR2 is located at the Xq28 locus, and it encodes arginine vasopressin receptor 2 (AVPR2). Mutations in AVPR2 have been associated with X-linked NDI. AQP2 is located at the 12q13.12 locus, and it encodes the water transporter aquaporin-2. Mutations in AQP2 result in autosomal NDI. Here, we describe a male infant with a novel AVPR2 variant who was referred to our hospital due to delayed head control.

Case Report

A 5-mo-old boy was referred to the Department of Child Neurology in Okayama University Hospital (Okayama, Japan) due to delayed head control. He was born to non-consanguineous parents. His birth weight was 3630 g (+1.39 SD) and birth length was 50.8 cm (+0.86 SD). He showed no signs of asphyxia. A recurrent fever of unknown origin was observed beginning at 2 mo of age. Poor weight gain was observed at 3 mo of age. Notably, he did not take any medications.

At his first visit to our hospital, the patient’s length was 62.6 cm (–1.55 SD) and weight was 6190 g (–1.82 SD). His heart rate was 128 bpm, and his body temperature was 36.8°C. He was dehydrated, and his head control was incomplete. Laboratory examination revealed that his sodium level and serum osmolarity were elevated, and his urinary osmolarity and specific gravity were noticeably low. Head magnetic resonance imaging scanning revealed that high posterior lobe intensity was absent.
in T1-weighted images. However, the pituitary stalk was intact, and no other abnormalities were identified. These findings were indicative of diabetes insipidus. Water deprivation test with vasopressin challenge test resulted in no urine osmolarity increase and no urine volume decrease. Moreover, the AVP level at admission was extremely high (114 pg/mL; reference range: 0.0–4.2 pg/mL). Based on these findings, we diagnosed the patient with congenital NDI. His urine volume decreased, his weight increased, and head control was achieved after initiation of hydrochlorothiazide.

Assessment of the patient’s family history revealed that his mother had polydipsia and polyuria from infancy until the present day (Fig. 1A). She also had increased levels of sodium of unknown origin during each of her three pregnancies. Furthermore, the patient’s maternal grandfather, maternal great-grandfather, and maternal aunt had histories of polyuria and polydipsia. We suspected that our patient had familial NDI and, thus, conducted a genetic analysis.

**Mutational Analysis**

Genetic analysis was approved by the ethical committee of Okayama University Hospital and conducted in accordance with the 1975 Declaration of Helsinki and subsequent amendments. Informed consent for genetic analysis was obtained from the patient’s mother.

We identified a hemizygous four-base pair duplication in exon 3 of AVPR2 (c.990_993 dup CAGC, Fig. 1B). This four-base pair duplication results in early termination (p.Val332Gln Fs26X). This variant was previously unreported, and it is not present in the 1000 Genomes Project databases (http://www.internationalgenome.org).
that reach the plasma membrane, but result in incomplete AVP binding. Class V mutations cause mis-sorting to incorrect cellular compartments. A duplication of four bases in exon 3 of AVPR2 causes early termination. However, this is the final exon, which might avoid nonsense-mediated RNA decay (3). Residue 332 is located in an intracellular domain that does not bind to AVP. Therefore, the variant in our patient may lose its function via a non-class I mechanism. Functional analysis is needed to clarify the pathophysiology of this variant.

Congenital NDI is generally identified by the presence of DI symptoms like polyuria, polydipsia, fever, and poor weight gain (2). The patient whose case is reported here was referred to the department of Child Neurology in our hospital because of delayed head control, poor weight gain, and fever. After treating his dehydration, he gained head control and his development proceeded normally. We consider delayed motor development in infancy to be an important clinical symptom of congenital NDI.

The patient’s mother had a heterozygous AVPR2 variant. She demonstrated a high sodium level during each of her pregnancies, but her sodium levels were normal before pregnancy and after birth for all children. In gestational DI, urinary volume increases during pregnancy because of increased clearance of AVP by placental vasopressinase, a cystine aminopeptidase that strongly degrades AVP (4). Although inactivation of the X chromosome was not analyzed in the patient’s mother, AVP resistance and degradation by vasopressinase, might have contributed to her high sodium levels during pregnancy.

In conclusion, we identified a novel AVPR2 variant in a patient with familial congenital NDI. Common symptoms of DI were observed, including polyuria, polydipsia, poor weight gain, and fever, and delayed motor development was also observed during infancy.

Conflict of Interest: The authors declare that they have no conflict of interest.

Discussion

We identified a novel AVPR2 variant in a patient with familial congenital NDI. Duplication variants are relatively rare among X-linked NDI patients: insertion AVPR2 variants were found in only about 5% of large cohort (1) and Japanese patients (2).

There are five classes of loss of function mutations in AVPR2 (1). Class I mutations result in unstable mRNA, which undergoes nonsense-mediated RNA decay. Class II mutations cause misfolding of receptors, and these mutant receptors remain in the endoplasmic reticulum. Class III mutations also cause receptor misfolding. Class III receptors reach the plasma membrane and react with AVP, but subsequent interactions with G proteins and cAMP production are impaired. Class IV mutations also result in misfolded receptors

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