Identification of a novel GRIN2D variant in a neonate with intractable epileptic encephalopathy—a case report

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Abstract

Background: N-methyl-D-aspartate (NMDA) receptors are ligand-gated ion channels that mediate excitatory synaptic transmission in the central nervous system. The functional NMDA receptors are heterotetramers consisting mainly of two GluN1 and two GluN2 subunits. GluN2 is encoded by the GRIN2D gene. A few case series have shown that GRIN2D variants are linked to developmental and epileptic encephalopathy. In this article, we report a novel GRIN2D variant, namely c.2021C > A (p.T674K) in a neonate with intractable epileptic encephalopathy.

Case presentation: A 12-day-old boy who had stiffness of the lower and upper extremities since birth was transferred from a local hospital to our department. On admission, the patient presented with head tilting backwards, staring, apnea and hypertonia of limbs. Video electroencephalogram showed continuous, generalized or multi-focal spike-wave and spike-and-slow wave discharges and hypsarrhythmia. A treatment regimen composed of phenobarbital, midazolam, levetiracetam and clonazepam was administered, which however led to only partial control of the seizure. Whole-exome sequencing identified c.2021C > A (p.T674K) in GRIN2D in the patient while such a mutation was not detected in the parents. The patient was hospitalized for 1 month and died of sudden cardio-respiratory arrest 2 weeks after discharge.

Conclusions: A novel variant of GRIN2D was identified in a neonate with epileptic encephalopathy. Epilepsy associated with this GRIN2D mutation is refractory to conventional anti-epileptic medications.

Keywords: case report, NMDA receptors, GRIN2D variants, de novo mutation, epileptic encephalopathy, anti-epileptic medications

Background

N-methyl-D-aspartate (NMDA) receptors are ligand-gated ion channels that are expressed throughout the central nervous system and mediate excitatory synaptic transmission [1]. To date, seven subunits of NMDA receptors, i.e., the GluN1 subunit, four distinct GluN2 subunits (GluN2A, GluN2B, GluN2C and GluN2D encoded by four different genes), and two GluN3 subunits (GluN3A and GluN3B) have been identified [2]. The functional NMDA receptors are heterotetramers consisting mainly of two GluN1 and two GluN2 subunits [3]. GluN2 is encoded by the GRIN2D gene [1]. A few case series have shown that GRIN2D variants are linked to developmental and epileptic encephalopathy [4–6]. In this study, we describe a novel de novo mutation in GRIN2D in a neonate with intractable epileptic encephalopathy.

Case presentation

A 12-day-old boy presenting with stiffness in the lower and upper extremities since birth was transferred from a local hospital to our department. He was born at full-term and had a normal delivery. The family history was unremarkable for neurological or seizure disorders. On physical examination, the patient was found to have hypertonia of the limbs, and repetitive eye deviation to the left side. His head control was limited, and he had a Restaurants of consciousness and frequent seizures. A treatment regimen composed of phenobarbital, midazolam, levetiracetam and clonazepam was administered, which however led to only partial control of the seizure. Whole-exome sequencing identified c.2021C > A (p.T674K) in GRIN2D in the patient while such a mutation was not detected in the parents. The patient was hospitalized for 1 month and died of sudden cardio-respiratory arrest 2 weeks after discharge.

Conclusions: A novel variant of GRIN2D was identified in a neonate with epileptic encephalopathy. Epilepsy associated with this GRIN2D mutation is refractory to conventional anti-epileptic medications.

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term without asphyxia to a G2P1 mother, with a birth weight of 3,600 g and an Apgar score of 9 and 10 at 1 and 5 minutes, respectively. Placenta, umbilical cord and amniotic fluid appeared normal. The parents were unrelated and healthy, and had no family history of epilepsy. On admission, the patient presented with head tilting backwards, staring, apnea and hypertonia in upper and lower limbs. He had a modified Ashworth scale score of 4 according to following findings: clenched fists with fingers fully flexed into the palm; flexed upper limbs to the chest; tonic extended lower limbs with flexed toes; and absence of sucking reflex, rooting reflex and Moro reflex. His temperature was 37 °C, heart rate 100/min, blood pressure 86/50 mmHg and SpO2 60%.

Blood, cerebrospinal fluid (CSF), urine and stool testing results were unremarkable. No growth was reported in blood and CSF culture. Cardiac enzyme profiling, electrolyte analyses, blood amino acid and acylcarnitine spectrum analyses for inherited metabolic diseases, and comprehensive panel of urine organic acids were unrevealing. Thyroid, liver and renal functions were normal. Cytomegalovirus and herpes virus, and their respective antibodies were negative as determined by PCR and ELISA, respectively. Chest radiography, electrocardiography, echocardiography and head MRI did not identify abnormalities. The automated auditory brainstem response was normal. Video electroencephalogram (EEG) was recorded and showed intermittent or continuous discharges (hypsarrhythmia) of generalized or multifocal spike-waves and spike-and-slow waves (Fig. 1), in line with epileptic encephalopathy.

The neonate was intubated and ventilated immediately after admission, and nasogastric feeding instituted. Phenobarbital, diazepam and vitamin B6 were initiated, which however did not show effect in seizure control (over 10 episodes/day). Diazepam was discontinued and a bolus intravenous injection of midazolam (0.15 mg/kg) was given followed by slower maintenance (0.2 mg/kg/h), which led to seizure reduction to ~ 10 episodes/day. On day 5 of admission, levetiracetam and clonazepam were added to the regimen, which significantly reduced seizure severity and frequency (5–6 episodes/day). Midazolam was tapered off in 2 weeks. Eighteen days after admission, the patient was extubated. Consent to taking a genetic test was obtained from the parents, and blood was drawn for whole-exome sequencing which was performed by Beijing Fulgent Technologies Inc. (Beijing, China). A heterozygous missense mutation, i.e., c.2021C > A (p. Thr674Lys) in GRIN2D in the patient was detected while such a mutation was not found in the parents (Fig. 2). The patient was discharged 1 month after hospitalization with continuous nasogastric administration of levetiracetam and clonazepam at home. Two weeks after discharge, the patient died of sudden cardio-respiratory arrest.
Discussion and Conclusions

To date, ten GRIN2D variants found in a total of 13 patients with developmental and epileptic encephalopathy have been described in the literature, which include Val667Ile, Met681Ile, Ser694Arg, Asp449Asn, Ser573Phe, Leu670Phe, Ala675Thr, Ala678Asp, Ser1271Leu and Arg1313Trp [4–6]. Clinical features, neurological context, treatment and outcomes of these patients have been summarized [4–6]. In this study, we identified a de novo missense mutation, i.e., Thr674Lys in GRIN2D, which is novel. According to the criteria for classifying pathogenic variants established by the American College of Medical Genetics and Genomics [7], the case we present here renders strong evidence of pathogenicity for this variant since (1) Thr674Lys was detected in the patient but not in the parents, (2) both maternity and paternity were confirmed, and (3) the patient had the disease without family history.

In the largest case series of 8 patients reported thus far, 7 received multiple anti-epileptic medications including clonazepam, levetiracetam, topiramate, vigabatrin and valproate, but only 2 completely responded to the treatment while 3 had no response and 2 achieved mild amelioration [4]. Li et al. studied 2 patients with Val667Ile and reported that Val667Ile, Leu670Phe, Ala675Thr and Ala678Asp had enhanced while Ser1271Leu and Arg1313Trp had reduced receptor activities [4, 6]. Of interest, based on the finding that Val667Ile is a gain-of-function mutation, Li et al. conducted a therapeutic trial using the NMDA receptor antagonist memantine to treat two patients with Val667Ile who were refractory to conventional antiepileptic medications, and reported mild to moderate improvement in seizure in both cases after memantine treatment [6]. In the future, functional testing of novel GRIN2D variants, which differentiates gain-of-function mutations from loss-of-function ones, may lead to more effective management of epilepsy by tailoring medical treatment to the individual characteristics of each variant, e.g., use of NMDA receptor antagonists for the management of epilepsy caused by GRIN2D gain-of-function mutations.

In conclusion, we identified a novel GRIN2D variant in a neonate with intractable epileptic encephalopathy. Our data together with previously reported findings suggest that epilepsy caused by GRIN2D mutations is predominantly refractory to conventional anti-epileptic therapies. Functional characterization of mutants might provide guidance for personalized medicine for epilepsy associated with GRIN2D mutations.
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Authors' contributions
JJ and LL drafted the manuscript. JJ, LL, MS, JF, LM1, YZ, CJ and LM2 participated in the acquisition, analysis, and interpretation of data. LM2 contributed to the conception of the study and substantively revised the manuscript. All authors read the final version of the manuscript and approved it for publication.

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Availability of data and materials
Data related to the case are presented in this article.

Ethics approval and consent to participate
As this is a case report, ethics approval is not required as per our institution’s policy. Informed consent was obtained from the parents of the patient.

Consent for publication
Written, informed consent was obtained from the parents of the patient for publication of this case report.

Competing interests
The authors declare that they have no competing interests.

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