

KCNQ2 encephalopathy manifesting with Rett-like features

A follow-up into adulthood

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Neurol Genet 2020;6:e510. doi:10.1212/NXG.0000000000000510

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Kv7.2 (KCNQ2) channel dysfunctions cause a rare form of neonatal and infantile epileptic and developmental encephalopathy (MIM 613720).¹ Unlike early clinical features and the epilepsy phenotype of KCNQ2 encephalopathy, data about the long-term developmental outcome are lacking. We followed up for over 3 decades a girl with this disease.

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▶ Video

The patient, who is aged 30 years and the only child of healthy unrelated parents, presented with pleiomorphic epileptic seizures (clonic, tonic asymmetric, and focal, with cyanosis) on the second day of life. These seizures were unresponsive to IV benzodiazepines and pyridoxine. EEG detected 2 independent foci in both frontotemporal regions. After a stormy onset, the combination of phenobarbital and phenytoin led to seizure improvement within the third month of life.

Neurologic examination during the first months of life detected minor facial dysmorphisms (figure), hypokinesia, and diffuse hypotonia. In the following years, progressive developmental delay, subcontinuous Rett-like hand stereotypies (flapping, washing, and mouthing) with reduced functional use of both hands, bradykinesia, hypomimia, severe dystonic quadriparesis, and kyphoscoliosis became obvious. She was never able to walk and had been using a wheelchair since age 7 years. At this stage, the girl showed profound intellectual disability (Griffiths Mental Development Scales: DQ 30) with absent language, hypoactivity, and severely limited social interaction. During late childhood and adolescence, epilepsy showed a relapsing-remitting course with focal clonic or generalized seizures and prolonged episodes of staring and unresponsiveness. Interictal EEG alterations included plurifocal spike and wave abnormalities predominantly in the anterior regions and a photoparoxysmal response at medium and high frequencies (figure, G). Between the ages of 17 and 20 years, seizures were well controlled with valproate and lamotrigine. After age 20 years, the patient remained seizure free with a stable clinical condition (video 1).

A brain MRI at age 4 years showed prevalent frontoinsular atrophy (figure, D–F). CSF examination at age 7 years detected normal glycorrachia, lactate, amino acids, biogenic amines, and pterins.

A next-generation sequencing panel that included 140 genes involved in genetic epilepsies revealed the de novo heterozygous KCNQ2 variant c.629G>A (p.Arg3894Gln), which was not present in the Human Gene Mutation Database (portal.biobase-international.com/). Its pathogenetic role is supported by previously published missense variants affecting the same amino acid in the S4 voltage sensor segment of the protein in patients with epileptic and developmental encephalopathies (c.628C>T; p.Arg210Cys and c.629G>A; p.Arg210His) and

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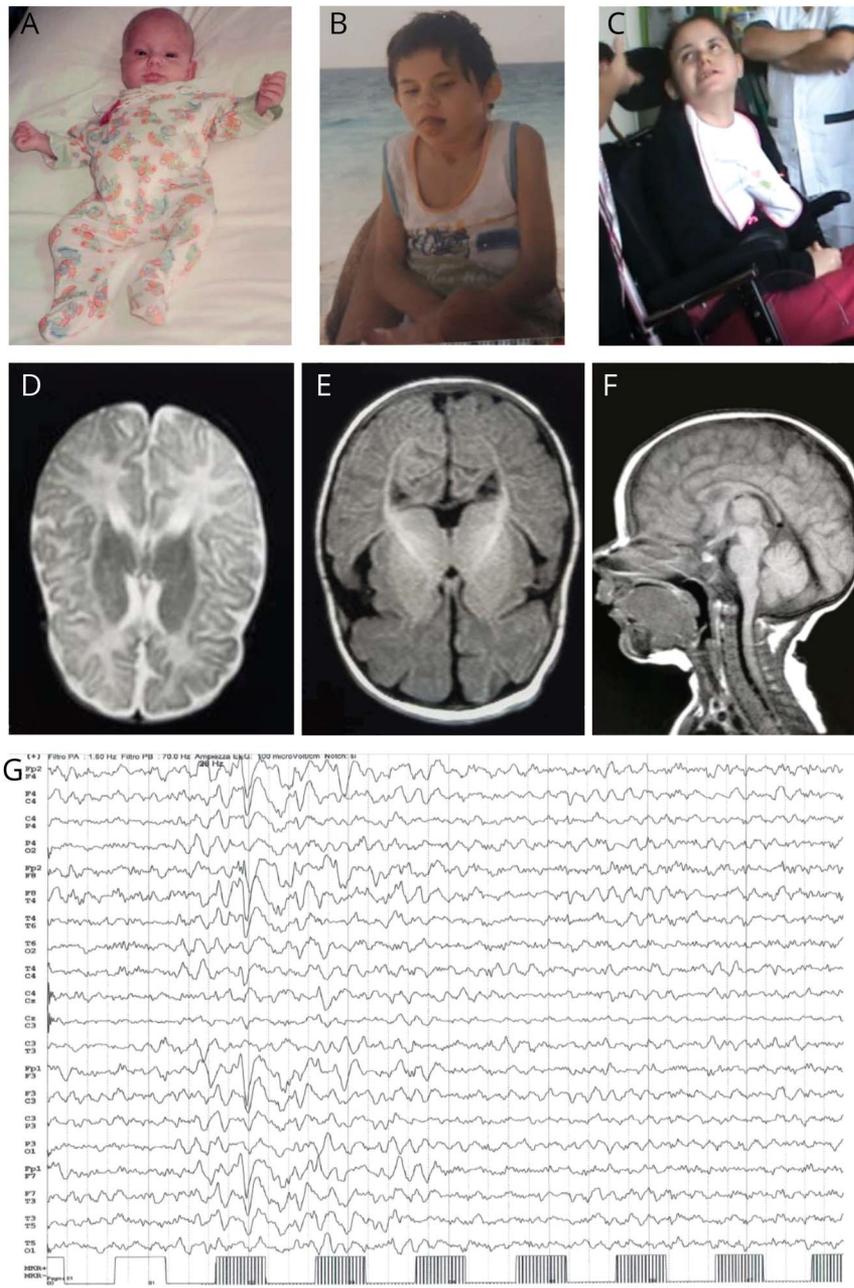
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Go to Neurology.org/NG for full disclosures. Funding information is provided at the end of the article.

The Article Processing Charge was funded by the authors.

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Figure Clinical, neuroradiologic, and EEG features of the reported patient



(A, B, and C) Minor facial dysmorphisms at age 5 months, 7 years, and 30 years (prominent ears and chin, deep-set eyes, and wide mouth with protruding tongue). (D, E, and F) Brain MRI at age 4 years showing frontotemporal cortical atrophy. (G) EEG at age 28 years showing a photoparoxysmal response at 28 Hz.

by multistate structural modeling studies that have demonstrated the involvement of the arginine at position 210 in the S4 segment in the stabilization of the protein through complex electrostatic interactions.¹⁻³ Although there were no therapeutic consequences, the molecular genetic diagnosis had a positive psychological impact on the parents.

The clinical history of our patient highlighted that motor and behavioral abnormalities, rather than epilepsy, characterize the adult phenotype of *KCNQ2* encephalopathy. The clinical follow-up of our patient showed the progressive emergence of a behavioral pattern resembling that of Rett syndrome. A

Rett-like presentation was previously reported in 8 patients with *KCNQ2* encephalopathy.^{4,5} Two of these patients carried the c.637C>T (p. Arg213Trp) variant, which involves the same region as the c.628C>T variant. One of them presented with neonatal seizures, global developmental delay, a lack of speech, and hand stereotypies.^{4,5}

Some parkinsonian symptoms, such as bradykinesia, hypomimia, hypoactivity, and dystonia (video 1), have previously been reported in classic *MeCP2*-related Rett syndrome and in other early-onset epileptic and developmental encephalopathies,⁶ suggesting that several genetically determined

neurodevelopmental disorders share a common adult outcome. However, unlike other diseases (including Kleefstra syndrome, *MECP2* mutations in males, and *CDKLS* encephalopathy in females), our patient did not experience a regression in cognitive or motor performance in adulthood.⁶

The role of Kv7.2 dysfunction in the expression of abnormal GABAergic parvalbumin-positive interneurons and their dendritic arborization within corticostriatal and nigrostriatal dopaminergic transmission (an impairment also described in *Mecp2*-null mice) could suggest a possible link between Rett syndrome, *KCNQ2* encephalopathy, and other neuropsychiatric disorders in terms of the pathophysiology of hypokinetic movement disorders, dystonia, and hand stereotypies.⁷ Recent experimental data on *KCNQ2*^{+/-} mice established a link between loss-of-function Kv7.2 mutations and behavioral disorders that was characterized by increased repetitive and compulsive behaviors, reduced social interest, and impaired social interaction mechanisms.⁷

From a clinical perspective, the present case highlights the importance of diagnostic reevaluation in adults with early-onset chronic neurologic conditions that were previously labeled according to a syndromic, rather than etiologic, diagnostic classification. In the absence of therapeutic consequences, the prognostic and psychological implications of a genetic diagnosis should not be underestimated.

Study funding

No targeted funding reported.

Disclosure

The authors have no disclosures to declare. Go to [Neurology.org/NG](https://www.neurology.org/NG) for full disclosure.

Publication history

Received by *Neurology: Genetics* April 16, 2020. Accepted in final form July 20, 2020.

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Filippo Manti, MD, PhD	Child Neurology and Psychiatry Unit, Department of Human Neuroscience, Sapienza University of Rome, Italy	Contributed to the design and conceptualization of the study and to the drafting of the manuscript
Maria Teresa Giannini, PT	Child Neurology and Psychiatry Unit, Department of Human Neuroscience, Sapienza University of Rome, Italy	Contributed to the drafting of the manuscript
Renzo Guerrini, MD, PhD, FRCP	Neuroscience and Neurogenetics Excellence Centre, Anna Meyer Children's Hospital, Florence	Revised the manuscript as senior author
Vincenzo Leuzzi, MD	Child Neurology and Psychiatry Unit, Department of Human Neuroscience, Sapienza University of Rome, Italy	Revised the manuscript as senior author

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Neurol Genet 2020;6;

DOI 10.1212/NXG.0000000000000510

This information is current as of September 8, 2020

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A Follow-up Into Adulthood

Neurol Genet 2021;7:e604. doi:10.1212/NXG.0000000000000604

In the Clinical/Scientific Note “KCNQ2 encephalopathy manifesting with Rett-like features: A follow-up into adulthood” by Mastrangelo et al.,¹ the first sentence of the fifth paragraph should read, “A next-generation sequencing panel that included 140 genes involved in genetic epilepsies revealed the de novo heterozygous *KCNQ2* variant c.629 G > C (p.Arg210Pro), which was not present in the Human Gene Mutation Database (portal.biobase-international.com/).” The authors regret the error.

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1. Mastrangelo M, Manti F, Giannini MT, Guerrini R, Leuzzi V. KCNQ2 encephalopathy manifesting with Rett-like features: a follow-up into adulthood. *Neurol Genet*. 2020;6(5):e510.