

Prenatal diagnosis of a nonsense mutation in the L1CAM gene resulting in congenital hydrocephalus: A case report and literature review

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Abstract. Congenital hydrocephalus is frequently caused by mutations in the L1 cell adhesion molecule (L1CAM) gene. The purpose of the present study was to identify possible causes of fetal hydrocephalus in a Chinese family. The samples from the parents and the hydrocephalic fetus were collected. Whole-exome sequencing and in-depth mutation analysis were performed. The identified variant, c.1267C>T.(p.Q423X), is situated on exon 11 of L1CAM gene (chromosome X:153134975). The fetus was confirmed to be hemizygous for the nonsense mutation and the mother was a heterozygous carrier. The mutation turns a glutamine into a premature stop codon at amino acid position 423. In conclusion, in the present study, a nonsense mutation in the L1CAM gene was identified during the prenatal diagnosis of a congenital hydrocephalic fetus from a Chinese family. The diagnosis highlighted the necessity of genetic screening for prenatal diagnosis.

Introduction

The neural L1 cell adhesion molecule (L1CAM) is a member of the immunoglobulin superfamily 1, constituted by six immunoglobulin-like domains, five fibronectin-III-like domains, a single pass transmembrane domain and a short cytoplasmic domain (1). It produces various molecular and cellular effects, which are key to nervous system development and are involved in neural migration and differentiation, as well as memory and learning processes (2-5).

The L1CAM gene is the known causative gene of L1 syndrome. It is close to the telomere on the long arm of the X chromosome and contains 28 exons (6). Mutation of the L1CAM gene may lead to four different X-linked neurological conditions, including hydrocephalus, mental retardation, spasticity of lower limbs and flexion deformity of the thumbs (7,8). The phenotype varies greatly from mild to severe, largely depending on the specific affected domains of the L1CAM protein. In detail, the mutations located on the extracellular domains generally cause severe conditions, while the pathologies associated with dysfunction of the cytoplasmic domains tend to be milder (9-11).

Hydrocephalus is defined as distension of the ventricular system of the brain owing to the inadequate passage of cerebrospinal fluid from the position of production within the cerebral ventricles to its region of absorption into the systemic circulation (12). Congenital hydrocephalus is one of the most common characteristics of L1 syndrome. It is most frequently caused by mutations in the L1CAM gene, which are responsible for 5-10% of male cases of isolated congenital hydrocephalus (13,14). Furthermore, the prognosis for this condition is poor.

The present study reported on a nonsense mutation in the L1CAM gene causing congenital hydrocephaly in a fetus from a Chinese family.

Case report

Case study. A 30-year-old healthy multipara visited The Second Affiliated Hospital and Yuying Children's Hospital of Wenzhou Medical University (Wenzhou, China) at 23 weeks of gestation on August 2018. She had already given birth to one healthy female infant and experienced one voluntary pregnancy termination at another hospital due to fetal hydrocephalus and the aborted male fetus was after the first girl was born. As indicated in Fig. 1, the familial pedigree suggested X-linked recessive hydrocephalus because the first girl was healthy, while both of aborted male fetuses presented with congenital hydrocephalus. As presented in Fig. 2, the ultrasound scan indicated progressive lateral ventricular dilatation, indicating severe fetal hydrocephalus. The maternal patient requested to terminate the pregnancy and provided written informed consent.

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To identify disease-causing factors, blood samples from the parents and umbilical cord blood of the aborted fetus were collected at 24 weeks of gestation. Preliminary karyotype analysis and single-nucleotide polymorphism array did not provide any evidence (15). Whole-exome sequencing (WES) and in-depth mutation analysis were then used to analyze the samples from the parents and umbilical cord blood (16). All underlying pathogenic mutants were confirmed by Sanger sequencing (6). Potential mutants were called by Genome Analysis Toolkit (GATK v3.7; <https://software.broadinstitute.org/gatk/download/>) (17) and the variants were filtered by gnomAD database (<https://gnomad.broadinstitute.org>; Exome Aggregation Consortium; ExAC Version 0.3.1) (17) and annotated by ANNOVAR (<https://annovar.openbioinformatics.org/en>; 2018 Mar 25 version) (17).

Literature search. Using PubMed (<https://pubmed.ncbi.nlm.nih.gov/>) as the primary search database, the literature was examined for nonsense mutations in the LICAM gene. PubMed searches of the references from database were performed using the key terms 'prenatal', 'hydrocephalus' and 'nonsense mutation'. Inclusion criteria were as follows: Fetal ventriculomegaly >15 mm; singleton pregnancies; normal karyotype and microarray; fetal ultrasound excluding additional central nervous system (CNS); or extra-CNS malformations. Cases with aqueductal stenosis or progressive enlargement of ventricles were excluded. A total of 11 publications related to the topic published between 1989 and October 25, 2020 were retrieved and their features are summarized in Table I. To date, 30 different nonsense mutations in the LICAM gene have been reported (7,8,16,18-27).

Results. The proband fetus was confirmed to be hemizygous for a mutant in the LICAM gene, NM_001278116.1:c.1267C>T:p.(Q423X), whilst the mother was identified to be a heterozygote and the father was healthy (Fig. 3). The nonsense mutation is located on exon 11 of the LICAM gene [chromosome (chr)X:153134975] and computational analysis [Sorting Intolerant From Tolerant (SIFT) (<https://genetics.bwh.harvard.edu/pph2>) (28) and PolyPhen-2 (<https://sift.jcvi.org/>) (29)] revealed that exon 11 is involved in encoding the fibronectin type-III of LICAM protein. According to the gnomAD database (<https://gnomad.broadinstitute.org>; Exome Aggregation Consortium; ExAC Version 0.3.1) (17), this mutation had not been previously reported among Asian populations. Based on the Online Mendelian Inheritance in Man database (OMIM; <https://www.omim.org>), disease-associated mutations in the LICAM gene were determined to be associated with X-linked hydrocephalus and the characteristics matched those of the proband. The identified mutation was X-linked recessive inheritance and pathogenic.

Discussion

Hydrocephalus, the abnormal accumulation of intracranial cerebrospinal fluid, is a common congenital malformation in fetuses. It occurs in 0.4-0.8% of newborns (14). The specific causes of this disease have remained to be fully identified. However, a number of ideas have been put forward. In a certain proportion of patients with genetic disorders,

congenital hydrocephalus manifests (13). It has been reported that X-linked hydrocephalus (XLH), which is caused by mutations in the LICAM gene, contributes to 5% of all cases of congenital hydrocephalus (10,14).

To date, >270 different pathogenetic mutations in the LICAM gene have been identified, including frameshifts, nonsense, missense mutations and alterations in splice-site junction (13). Certain studies have explored the associations between specific mutations in the LICAM gene and the severity of syndromes. Mutations leading to extracellular protein truncation may produce severe pathogenesises, whereas mutations in cytoplasmic domains tend to lead to milder manifestations (7,9-11).

The criteria for identifying a high-risk mutation in the LICAM gene are family history and the clinical LI syndromes, including stenosis of the aqueduct of Sylvius, mental retardation, aphasia, spastic paraplegia, adducted thumbs, X-linked complicated hereditary spastic paraplegia type 1 and X-linked agenesis of the corpus callosum (7,8,10,13). In the present study, in view of that the patient had experienced a previous pregnancy with fetal hydrocephalus and the current fetal ultrasound revealed hydrocephalus, genetic causes were suspected. A nonsense mutation, NM_001278116.1:c.1267C>T:p.(Q423X) in exon 11 of the LICAM gene (chrX:153134975) was identified in the Chinese family. The mutant converts a glutamine into a premature stop codon on amino acid position 423, which in turn leads to extracellular protein truncation that abrogates the actions of LICAM in its intracellular signaling pathways (7,18,19). However, there was no history of hydrocephalus in the mother's family. Nonsense-mediated mRNA decay (NMD) is an RNA surveillance mechanism, which may be used to detect mRNAs that harbor premature termination codons and initiate the degradation of the truncated or potentially harmful proteins. In the present case, it may have been suitable to perform NMD analysis of the nonsense mutation. However, due to improper preservation of specimens, this was not performed. In a future study, it may even be possible to trigger the RNA surveillance mechanism artificially to observe whether the harmful effects resulting from the nonsense mutation may be alleviated by NMD.

Since only one case has so far been reported, it is impossible to clarify the exact mechanisms of how a mutation at this site causes disease in individuals. This is a limitation of this study. However, it is possible to estimate the pathogenicity of a mutation with some available online tools, e.g., MutationTaster. Specifically, the pathogenesis observed to result from nonsense mutations on other sites in the LICAM gene may provide clues. To date, 30 different nonsense mutations in the LICAM gene have been reported (7,18,21-27). All of the mutations were unique to each family, with the exception that one mutation appeared in two different families (c.3496C>T) (7,23) and two different mutations were detected in one family (c.614C>A, c.615C>G) (7). All of these 30 mutations are considered capable of causing disease.

The characteristics of XLH include the enlargement of both third and lateral ventricles, and agenesis of the corpus callosum (7,8). Previous data revealed that 41% of patients with congenital hydrocephalus carried pathogenic mutations in the LICAM gene (26). Furthermore, the prognosis of these carriers

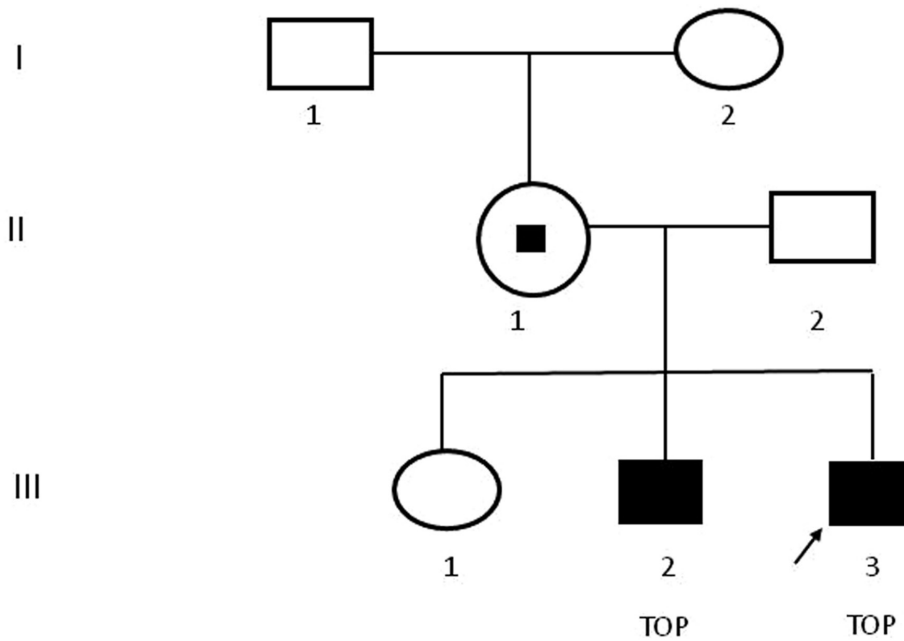


Figure 1. Pedigree chart. Females are indicated with circles and males with squares. The proband is indicated by an arrow. Filled squares indicate affected individuals and the small black square indicates mutation carrier status. TOP, termination of pregnancy.

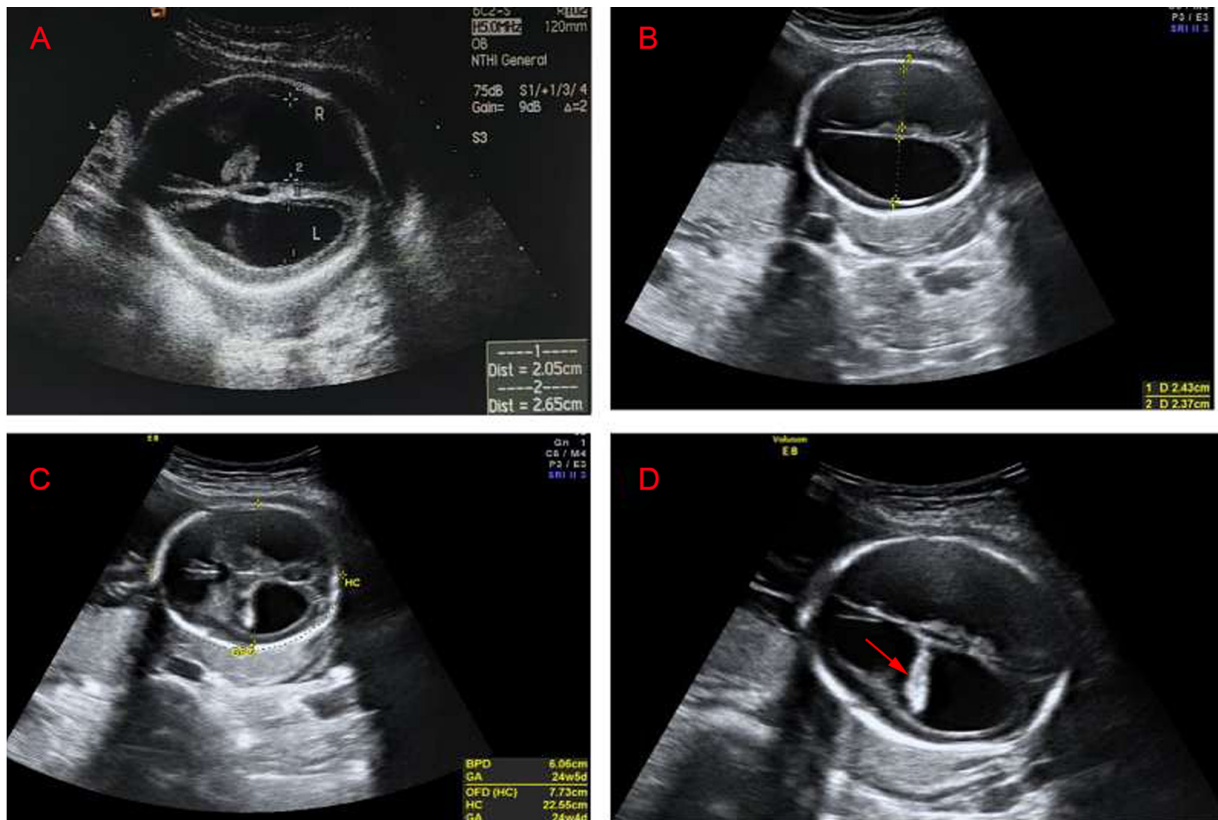


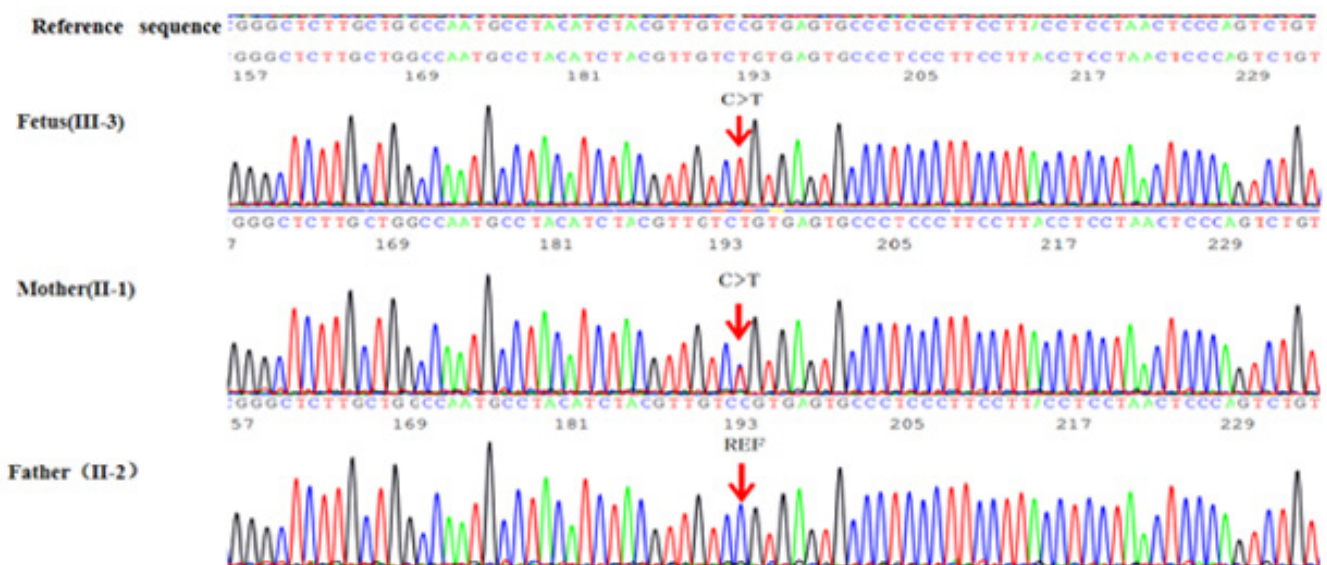
Figure 2. Transabdominal view of the fetal brain. (A) Ultrasound image of the first hydrocephalus fetus. The width of the left ventricle was 2.05 cm, the width of the right ventricle was 2.65 cm and the cerebral cortex became thinner. (B) Ultrasound image of the second hydrocephalus fetus. The width of the left ventricles was 2.43 cm, the width of the right ventricle was 2.37 cm and the cerebral cortex became thinner. (C) Ultrasound image of the second hydrocephalus fetus. The biparietal diameter of the fetus was 6.06 cm, the occipitofrontal diameter was 7.73 cm and the head circumference was 22.55 cm. (D) Ultrasound image of the second hydrocephalus fetus. Hyperechoic choroid hanging sign (arrow), the bilateral lateral ventricles were expanded and the cerebral cortex became thinner.

is exclusively worse. In certain cases, hydrocephaly in fetuses may cause stillbirth, neurodevelopmental delay in survivors

or death in early infancy (13). Facing such conditions, most parents choose to terminate their pregnancies. However, in other

Table I. Nonsense mutations in the L1 cell adhesion molecule gene.

Author (year)	Position in the gene (exon no.)	Nucleotide change	Consequence	Country	(Refs.)
Gu <i>et al</i> , 1996	08	c.828G>A	p.Trp276Ter	Germany	(21)
Gu <i>et al</i> , 1997	11	c.1318C>T	p.Gln440Ter	Germany	(22)
	23	c.3124C>T	p.Gln1042Ter		
Finckh <i>et al</i> , 2000	04	c.316C>T	p.Q106X	Germany	(7)
	11	c.1267C>T	p.Q423X		
	26	c.3496C>T	p.R1166X		
	02	c.79G>T	p.E27X		
	21	c.2746G>T	p.G916X		
	06	c.614C>A c.615C>G	p.S205X		
	13	c.1672C>T	p.R558X		
	14	c.1786G>T	p.E597X		
Okamoto <i>et al</i> , 2004	22	c.2974C>T	p.Gln992Ter	Japan	(8)
Kanemura <i>et al</i> , 2006	26	c.3496C>T	p.R1166X	Japan	(23)
Jackson <i>et al</i> , 2009	13	c.1674C>T	p.R558X	USA	(24)
Takenouchi <i>et al</i> , 2012	01	c.61C>T	p.Q21X	Japan	(25)
Adle-Biassette <i>et al</i> , 2013	06	c.621C>A	p.Tyr207X	France	(26)
	07	c.772C>T	p.gln258X		
	08	c.871C>T	p.gln291X		
	08	c.892C>T	p.gln298X		
	12	c.1380G>A	p.Trp460X		
	12	c.1453C>T	p.Arg485X		
	12	c.1453C>T	p.Arg485X		
	15	c.1882C>T	p.gln628X		
	16	c.1989C>G	p.Tyr663X		
	21	c.2785C>T	p.gln929X		
	23	c.3163G>T	p.gly1055X		
	23	c.3107G>A	p.Trp1036X		
Marín <i>et al</i> , 2015	26	c.3478C>T	P.Gln1160X	Spain	(27)
Ochando <i>et al</i> , 2016	18	c.2239C>T	P.Gln747X	Spain	(18)
Guo <i>et al</i> , 2020	21	c.2865C>A	p.Y955X	China	(16)

Figure 3. Gene sequencing results in L1 cell adhesion molecule for the fetus and the parents. The ref sequence was cited from Human GRCh37/hg19 (<https://ncbiinsights.ncbi.nlm.nih.gov/2020/11/12/human-grch37-hg19-refseq-annotation-update/>). The mutation site is indicated by arrows.

cases, terminating pregnancy was not a legal option given the gestational age. Intrauterine ventriculo-amniotic shunting is an option for managing such cases. Theoretically, such therapy is beneficial for preventing amniotic fluid reflux and cerebral injury. The efficacy of this treatment has been proven in animal experiments (30). However, attempts in the 1980s were generally abandoned due to the high prevalence of severe neurodevelopmental delay among survivors (31). Recently, a retrospective study including 44 fetuses with severe bilateral ventriculomegaly treated with ventriculo-amniotic shunting demonstrated the feasibility of this treatment for severe fetal cerebral ventriculomegaly. However, repeated shunting was required in nearly half of the cases and a high proportion of survivors have neurodevelopmental delay. It has been proposed that the possible beneficial role of ventriculo-amniotic shunting requires to be confirmed by randomized studies with a large sample size (32).

There are currently no effective therapies for curing genetic disabilities. The only available method is to prevent the birth of fetuses with diagnosed hydrocephalus and the reoccurrence of genetic hydrocephalus in future pregnancies. Comprehensive prenatal diagnostic measures have considerable significance in assisting clinicians in making informed decisions and providing guidance for families with LICAM gene mutants.

In addition, prenatal ultrasound may detect the presence of hydrocephalus, but its dependability is not optimal. Kanemura *et al* (23) demonstrated only 21.7% of cases of XLH were identifiable by ultrasound screening prior to 22 weeks of gestation. Of note, enlargement of the ventricles usually does not begin until 20 weeks of gestation and certain individuals may only develop this condition after live birth or not at all (33). On one hand, prenatal diagnosis of hydrocephalus cannot always be made as early as possible. On the other hand, false-negative results undoubtedly increase the clinical variability for diagnosing L1-associated disease (33). Accordingly, using DNA sequencing to identify the genetic causes of hydrocephalus is particularly important in families carrying LICAM gene mutations. It has been reported that the detection rate of LICAM gene mutations in patients without a family history of L1 syndrome is 9-50%, while among those with a family history of L1 syndrome, the probability of LICAM gene mutations is up to 50-80% (7,10,34).

In conclusion, the present study reported on a nonsense mutation, NM_001278116.1:c.1267C>T:p.(Q423X), in the LICAM gene as the probable cause of XLH in a Chinese family. According to the diagnostic experience in this case, prenatal ultrasound and DNA screening should be offered to high-risk families to assist them in deciding whether to continue a pregnancy (14). Determining which mutations are associated with fetal malformations may also improve counseling for future pregnancies.

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Availability of data and materials

The datasets generated and/or analyzed during the current study are available in the LOVD database repository (<https://databases.lovd.nl/shared/variants/0000796593#00025606>).

Authors' contributions

RW, HC, and XWu performed genetic diagnoses and drafted and critically revised the manuscript. RW and XWu Authenticated the raw data in this manuscript. RW, XWa, SH, and AX clinically diagnosed patients, collected and analysed samples and performed imaging tests. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

This study has been approved by the Ethics Committee/Institutional Review Board of the Second Affiliated Hospital and Yuying Children's Hospital of Wenzhou Medical University (Wenzhou, China). Written informed consent, according to the Declaration of Helsinki, were acquired from both parents of the fetus.

Patient consent for publication

All probands provided informed consent for genetic testing and publication of their data in this study.

Competing interests

The authors declare that they have no competing interests.

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