INTRODUCTION

Cases of distal 15q chromosomal duplication have been reported in the literature but are very uncommon. After the first case with a duplication of distal 15q was reported in 1974,1 approximately 100 cases have been documented, but with few de novo duplications as in the present case. Subsequent case studies showed that chromosome region 15q11-q13 was a hot spot for chromosomal duplications.

Abstract

Background: Terminal duplication on chromosome 15q is a rare chromosomal variation. Affected individuals show similar features such as growth dysplasia or the development of frontal bossing, body deformities, facial abnormalities, and genitourinary or cardiovascular disorders. However, it is not yet clear whether such 15q repeats lead to identifiable patterns of clinical abnormalities. Therefore, the purpose of this study was to analyze the prenatal diagnostic results and clinical manifestations of a fetus with 15q duplication and to summarize the literature.

Methods: The case was a fetus at 28 weeks of gestation. The risk of Down syndrome from second-trimester screening was 1/140. Prenatal ultrasound and amniocentesis were performed, and chromosomal microarray analysis (CMA) was used for genetic analysis.

Results: The fetus had abnormal clinical features, including intracardiac echogenic focus in the left ventricle, an aberrant right subclavian artery, and growth delay. The fetal chromosomal karyotype was 46,XX,15q?,12q?,21pstk+, and CMA revealed a 10.163 Mb duplication at 15q24.3-q25.3. The couple chose to terminate the pregnancy after careful consideration.

Conclusions: The combination and rational application of cytogenetics technology and molecular genetics technology such as CMA will open up the field of clinical application and provide useful genetic counseling for parents of fetuses carrying such chromosomal duplications.

KEYWORDS
15q duplication, abnormal ultrasound, chromosomal microarray analysis, genetic counseling, prenatal diagnosis
duplication. However, there are few de novo duplications in the region of 15q24-pter. Previous studies have described some cases of 15qter duplication characterized by postnatal or prenatal overgrowth, craniofacial and skeletal malformations, developmental delay, and genital abnormalities. The significant abnormalities in fetal growth and development, and the formation of congenital malformations are caused by the abnormal expression of genes located in the 15qter region such as LINGO-1, CSPG4, MTHFS, KIF7, CHD2, and IGF1R. Moreover, the range and severity of symptoms, and physical findings are closely related to the length and location of the duplicated region of chromosome 15q, and these can vary from case to case. For example, patients with 15q duplications exhibit some clinical phenotypes that are opposite to overgrowth, such as postnatal or prenatal growth restriction and developmental delay.

Here, we reported a fetus with a double de novo duplication of chromosome 15q24.3-q25.3, producing abnormal sonography findings. The duplication was detected by chromosomal microarray analysis (CMA), and we identified 19 potentially pathogenic genes including LINGO-1 and MTHFS using the DECIPHER genome browser (https://decipher.sanger.ac.uk). To the best of our knowledge, no case of partial tetrasomy of 15q24.3-q25.3 has been reported previously.

2 MATERIALS AND METHODS

2.1 Case report

A 28-year-old primigravid woman underwent amniocentesis for prenatal diagnosis at 28 weeks of gestation because second-trimester screening for Down syndrome indicated a high risk (1/140), calculated from abnormal maternal serum screening markers. The multiple of median (MoM) values of maternal serum screening markers were as follows (Table 1): 0.707 MoM for free beta-human chorionic gonadotropin (Free β-hCG), 0.503 MoM for unconjugated estriol (uE3), and 0.362 MoM for alpha-fetoprotein (AFP). Noninvasive prenatal testing was performed at 16 weeks of gestation (ie, second trimester), but the results showed a low risk of chromosomal aneuploidies. Fetal exfoliated cells in amniotic fluid were used for karyotyping and CMA. Color Doppler echocardiography at 28 weeks of gestation revealed an intracardiac echogenic focus in the left ventricle (Figure 1A) and an aberrant right subclavian artery (Figure 1B). Table 1 shows the results of systemic ultrasonography at 28 weeks of gestation. Fetal abdominal circumference, head circumference, humerus length, and weight were all low for gestational age (<10th centile). The fetus’s parents were not consanguineous and were healthy. The mother denied being exposed to teratogenic agents or irradiation, or using nicotine, alcohol, or caffeine during the pregnancy. No family history of genetic disease, congenital malformations, or diabetes mellitus was recorded. The study protocol was approved by the Ethics Committee of the First Hospital of Jilin University, and written informed consent was obtained from the couple. Informed consent for publication of this case has also been provided by the couple.

| Indicator                          | Values | Normal range |
|-----------------------------------|--------|--------------|
| Maternal serum screening results  |        |              |
| AFP (MoM)                         | 0.362  | 0.7-2.5      |
| Free β-hCG (MoM)                  | 0.707  | 0.25-2.0     |
| uE3 (MoM)                         | 0.503  | 0.5-2.0      |
| DS risk                           | 1/140  | <1/270       |
| Evaluation of amniotic fluid      |        |              |
| Maximum deepest vertical pocket (cm) | 2.91  | 2.8          |
| Amniotic fluid index (AFI) (cm)   | 8.53   | 5-24         |
| Fetal heart rate (FHR) (times per minute) | 140   | 110-160      |
| Ultrasound indicators of fetal size (28 wk) |        | (10-90th centiles) |
| Biparietal diameter (cm)          | 7.02   | 6.8-7.5      |
| Femur length (cm)                 | 5.26   | 5.0-5.5      |
| Abdominal circumference (cm)      | 20.78  | 22.6-25.2    |
| Head circumference (cm)           | 25.09  | 25.2-27.2    |
| Humerus length (mm)               | 43.9   | 45.3-51.7    |
| Fetal weight (g)                  | 963    | 995-1404     |

Abbreviations: AFP, alpha-fetoprotein; DS, Down syndrome; Free β-hCG, free beta-human chorionic gonadotropin; MoM, multiple of median; uE3, unconjugated estriol.
karyotyping. The karyotype description refers to the International System for Human Cytogenetic Nomenclature (ISCN 2013). Because of the abnormal karyotype of the fetus, the parents were recalled for further such tests after obtaining written informed consent.

2.3 | Chromosome microarray and data analysis

Ultrasound-guided amniocentesis was performed to extract about 10 mL of amniotic fluid for CMA. DNeasy Blood & Tissue kits (Qiagen GmBH) were used to extract genomic DNA according to the manufacturer’s instructions. A NanoDrop ND-2000 spectrophotometer (Thermo Fisher Scientific) was used to quantify the DNA. The potential copy number variations (CNVs) were detected using an Affymetrix CytoScan750K Array (Affymetrix). DNA processing included digestion, joining, breaking, marking, hybridization, staining, and scanning. Software of the chromosome analysis suite (ChAS) was used to analyze the data. The array data and genotype-phenotype correlations were analyzed by using the databases of Genomic Variants (http://dgv.tcag.ca/dgv/app/home; GRCh37/hg19), OMIM (https://omim.org), and DECIPHER (see above).

3 | RESULTS

Initially, the fetus was diagnosed with an abnormal karyotype of 46,XX,15q?,12q?,21pstk+ (Figure 2A) by routine cytogenetics for prenatal diagnosis. CMA revealed a 10.163 Mb duplication of 15q24.3-q25.3 at twice [15q24.3-q25.3 [15:77 456 021-87618593] × 4] (Figure 3), but no abnormalities were found on chromosome 12q. Conventional cytogenetics demonstrated that the mother had a chromosomal polymorphism of 21pstk+ (Figure 2B) but the father had a normal karyotype. Furthermore, the couple asked for a CMA study, and all results were normal. Thus, the chromosome 15q24.3-q25.3 duplication of the fetus detected by CMA was a de novo chromosomal variation. Finally, the couple chose to terminate the pregnancy after careful consideration because of the abnormal ultrasonography findings and CMA results.

**FIGURE 1** Prenatal ultrasound findings at 28 wk of gestation: A, Intracardiac echogenic focus in the left ventricle; B, Aberrant right subclavian artery

**FIGURE 2** A, Karyotype of the fetus identified by GTG banding technique. B, The mother’s karyotype

**FIGURE 3** A, Karyotype identified by GTG banding technique of the fetus with a 10.163 Mb duplication of 15q24.3-q25.3.
4 | DISCUSSION

Trisomy or tetrasomy for chromosome 15qter is very rare; about 30 patients have been described in the literature to date. Affected individuals exhibit similar clinical features, including cephalic or facial deformities, osteoarticular abnormalities, a subarachnoid space, and renal, urogenital or cardiovascular diseases. Here, we report another fetus with a nonmosaic tetrasomy at 15q24.3-q25.3 resulting from duplication of chromosome 15qter. The fetus exhibited abnormalities including an intracardiac echogenic focus in the left ventricle, an aberrant right subclavian artery, growth delay and low amniotic fluid index, and a double de novo duplication of 15q24.3-q25.3 identified by CMA. To illustrate genotype-phenotype correlations, the similar clinical features of individuals with 15q24-qter duplications are summarized in Table 2.

A total of 16 cases listed in Table 2 were pure 15q duplication, and which covers the 15q24.3-25.3 region. Among the 16 cases, there were 4 cases with tetrasomy for 15qter and 12 cases with trisomy. The detection methods are as follows: 2 cases detected by karyotype, 9 cases by karyotype combined with FISH, and 5 cases detected by CMA.

According to the literature, we inferred that the clinical features of 15q24-qter duplication are genetic disorders that may be caused by nonallelic homologous recombination between low-copy repeats in the region of chromosome 15q24-qter. The 15q24-q26 region is one of several hotspots reported, with a high density of chromosome-specific duplications. Rearrangements of this region have been implicated as a susceptibility factor for panic and phobic disorders with joint laxity. However, according to case reports, it is not yet clear whether the 15q repeats lead to identifiable patterns of clinical abnormalities. Zollino et al summarized 32 patients with 15q duplications and divided them into two groups: The first had duplication at 15q21-24qter, revealing normal prenatal growth and microcephaly; the second had duplication of 15q25-26qter, leading to macrocephaly, prenatal overgrowth, and craniosynostosis. Gutierrez-Franco Mde et al reported a case of trisomy of distal 15q with overgrowth and mental retardation. Another case report described a newborn infant with a de novo 15q24-q26 duplication and intrauterine overgrowth. However, ultrasonography indicated that the fetus in this case might have had growth delay, but there was no sign of head size deformity.

Mental retardation seems to be a common feature of patients with duplication of 15qter, because several genes related to brain development and function are involved in the 15q24.3-qter region. Schluth et al, Liehr et al, and Alakbarzade et al reported cases of clinical manifestations of mental retardation. Another common feature of individuals with duplication of 15qter is cardiac malformation, which has been found in about 50% of the cases. The types of cardiac malformations include...
| References          | Yip et al<sup>19</sup> | Blennow et al<sup>20</sup> | Abe et al<sup>21</sup> | Roggenbuck et al<sup>9</sup> | Genesio et al<sup>10</sup> | Schluth et al<sup>22</sup> |
|---------------------|------------------------|-----------------------------|-------------------------|-----------------------------|---------------------------|-----------------------------|
| Age/sex             | NR                     | caseA 2 y/M                 | caseB 3 y/F             | 2 mo/F                      | case1 died/F              | case2 4.5 y/F               | case3 21 mo/M              | 3 wk/F                      | 23 y/M                    |
| Duplicated region   | 15q21-qter             | 15q23-qter                 | 15q24-qter             | 15q24.3-q26.3              | 15q24-q26.3               | 15q24-q26.3                 | 15q21.3-q26.3              | 15q24.3-qter               |
| Facial dysmorphism  | +                      | +                           | +                       | NR                          | +                         | +                           | +                         | +                           |
| Malformation of finger or toe | +                   | +                           | +                       | NR                          | +                         | +                           | +                         | +                           |
| Muscular hypotonia  | +                      | NR                          | NR                      | NR                          | NR                        | NR                          | NR                        | +                           |
| Convulsive seizures or tremor | +                   | NR                          | NR                      | NR                          | NR                        | NR                          | NR                        | +                           |
| Osteoarticular abnormality | −                    | +                           | +                       | NR                          | −                         | −                           | −                         | +                           |
| Arachnodactyly      | NR                     | +                           | +                       | NR                          | NR                        | −                           | NR                        | +                           |
| Achromatopsia       | NR                     | +                           | −                       | NR                          | NR                        | NR                          | NR                        | NR                          |
| Sensorineural hearing defect | +                   | +                           | +                       | NR                          | NR                        | +                           | −                         | NR                          |
| Mental retardation  | +                      | +                           | +                       | NR                          | −                         | NR                          | −                         | NR                          |
| Postnatal overgrowth | −                     | +                           | +                       | NR                          | NR                        | +                           | −                         | NR                          |
| Developmental or growth delay | +                 | NR                          | NR                      | NR                          | +                         | −                           | +                         | NR                          |
| Cardiac Malformations | −                    | −                           | −                       | +                           | NR                        | +                           | +                         | NR                          |
| Nural tube defects  | −                      | −                           | −                       | −                           | +                         | −                           | −                         | −                           |
| Short neck          | −                      | −                           | −                       | −                           | NR                        | +                           | −                         | −                           |
| Genital abnormality | NR                     | −                           | −                       | NR                          | NR                        | +                           | −                         | NR                          |
| Brachycephaly       | −                      | −                           | −                       | NR                          | NR                        | +                           | −                         | −                           |
| Renal dysplasia     | NR                     | NR                          | NR                      | NR                          | NR                        | NR                          | +                         | +                           |
| Intrauterine overgrowth | NR                  | NR                          | NR                      | NR                          | NR                        | NR                          | NR                        | −                           |
| Single umbilical artery | NR                   | NR                          | NR                      | NR                          | NR                        | NR                          | NR                        | NR                          |

| References          | Liehr et al<sup>23</sup> | Gutierrez-Franco Mde et al<sup>24</sup> | Kim et al<sup>7</sup> | O’Connor et al<sup>25</sup> | Chen et al<sup>6</sup> | Ochando et al<sup>6</sup> | Alakbarzade et al<sup>26</sup> | Present case |
|---------------------|--------------------------|--------------------------------------------|-----------------------|--------------------------|------------------------|--------------------------|-------------------------------|---------------|
| Age/sex             | 11 y/F                   | 13 y/F                                    | Newborn/F              | Newborn/F                | Fetus/F                | Fetus/M                  | Pedigree                      | Fetus/F        |
| Duplicated region   | 15q24.1-qter             | 15q24-q26.3                               | 15q24-q26.3            | 15q24.2-q26.3            | 15q24.2-q26.2          | 15q24.3-q26.1            | 15q24.3-q25.1               | 15q24.3-q25.3 |
| Facial dysmorphism  | +                        | +                                         | +                      | +                        | NR                     | NR                      | −                            | −              |
| Malformation of finger or toe | +                      | +                                         | +                      | +                        | NR                     | NR                      | NR                            | −              |
| Muscular hypotonia  | +                        | −                                         | NR                     | NR                      | NR                     | NR                      | +                            | −              |
| Convulsive seizures or tremor | +                      | −                                         | NR                     | NR                      | NR                     | NR                      | NR                            | +              |
| Osteoarticular abnormality | +                    | +                                         | +                      | NR                      | NR                     | NR                      | NR                            | −              |
| Arachnodactyly      | NR                       | NR                                         | NR                     | NR                      | NR                     | NR                      | NR                            | NR             |
Wolf-Parkinson-White syndrome, mitral valve stenosis, Ebstein’s anomaly, mitral valve arcade, defects of the atrial and ventricular septum, an atrioventricular canal, subaortic stenosis, patent foramen ovale or ductus arteriosus, cardiomegaly, and an aberrant right subclavian artery (as in the present case). Here, we also found cardiac malformations manifested as an intracardiac echogenic focus in the left ventricle. Obviously, we could not evaluate whether the fetus had mental retardation, because this can only be observed after birth and ultrasonography can only indicate a structural abnormality.

ADAMTS13 (cytogenetic location: 15q25.2; OMIM #609199), a gene associated with the duplication of chromosome 15qter has been identified as a potential cause of cardiac and vessel malformations. Overexpression of the ADAMTS13 gene has also been thought to interfere with kidney function. Genesio et al. reported a female infant with a 15q21.3-q26.3 duplication and a horseshoe kidney, and Kim et al. reported a newborn with a 15q24-q26.3 duplication with hydronephrosis. However, no fetal kidney abnormality was found in our case. All the genes involved in this region are shown in Figure 4. The 15q24.3-q25.3 region contains 19 morbid genes involved in morbidity, namely LINGO1, CIB2, IREB2, CHRNA5, CHRNA3, MIR184, MTHFS, FAH, ARNT2, MESDC2, EFTUD1, RPS17, RPS17L1, AP3B2, HOMER2, WDR73, ALPK3, SLC28A1, and AGB1. The corresponding phenotype of these genes and explanations are summarized in Table 3. LINGO-1 and MTHFS have been shown to be related to the clinical manifestations.

Traditional chromosome banding and karyotyping have always been the gold standard of cytogenetics and have irreplaceable advantages. In the present case, the fetal karyotype was 46,XX,15q?,12q?,21pstk+. The mother was 46,XX,21pstk+, and the father was normal (46,XY). The fetal 21pstk+ chromosomal polymorphism was inherited from the mother, but this is considered to have no detrimental phenotypic effect. Furthermore, CMA results demonstrated a de novo duplication of 15q24.3-25.3. Hence, further prenatal counseling for the pregnant woman and her family could be given appropriately. However, the results were complicated; CMA works by detecting imbalances in DNA copy numbers, or CNVs. Therefore, based on our results and theoretical knowledge, we speculate that the 15q duplication fragment might have been a partially balanced translocation with 12q, which is probably why CMA could not detect a chromosome 12q abnormality. Clearly, the origin of any abnormal fetal chromosome 12q found by karyotyping needs to be verified by other techniques, such as fluorescence in situ hybridization (FISH). Further FISH verification can indeed identify the location of the duplication and verify the diagnostic results. However, there were no remaining samples after completing the karyotype and CMA, and the fetus had induced labor. This is the limitation of the present study.
FIGURE 4  The involving genes contained in the region of 15q24.3-25.3 (15:77456021-87618593). The figure is modified from the DECIPHER genome browser

TABLE 3  Morbid genes in the region of 15q24.3-25.3 and the associated phenotype

| Gene     | location | OMIM   | Explanation                                                                 | Phenotype                                      |
|----------|----------|--------|-----------------------------------------------------------------------------|------------------------------------------------|
| LINGO1   | 15q24.3  | 609791 | Leucine-Rich Repeat- And Ig Domain-Containing Nogo Receptor-Interacting Protein 1 | Mental retardation, autosomal recessive 64     |
| CIB2     | 15q25.1  | 605564 | Calcium- And Integrin-Binding Protein 2                                     | Deafness, autosomal recessive 48, Usher syndrome, type IJ |
| IREB2    | 15q25.1  | 147582 | Iron responsive element binding protein 2                                   | Neurodegeneration, early-onset, with choreoathetoid movements and microcytic anemia |
| CHRNA5   | 15q25.1  | 118505 | Cholinergic Receptor, Neuronal Nicotinic, Alpha Polypeptide 5                | Smoking as A Quantitative Trait Locus 3; SQTL3  |
| CHRNA3   | 15q25.1  | 118503 | Cholinergic Receptor, Neuronal Nicotinic, Alpha Polypeptide 3                | Smoking as A Quantitative Trait Locus 3; SQTL3  |
| MIR184   | 15q25.1  | 613146 | Micro Rna 184                                                               | EDICT syndrome                                 |
| MTHFS    | 15q25.1  | 604197 | 5,10-Methenyltetrahydrofolate Synthetase                                     | Neurodevelopmental disorder with microcephaly, epilepsy, and hypomyelination |
| FAH      | 15q25.1  | 613871 | Fumarylacetoacetate Hydrolase                                               | Tyrosinemia, type I                            |
| ARNT2    | 15q25.1  | 606036 | Aryl Hydrocarbon Receptor Nuclear Translocator 2                            | Webb-Dattani syndrome                          |
| MESDC2   | 15q25.1  | 607783 | Mesoderm Development Candidate Gene 2                                       | Osteogenesis imperfecta, type XX               |
| EFTUD1   | 15q25.2  | 617538 | Elongation Factor-Like Gtpase 1                                             | Shwachman-Diamond syndrome 2                   |
| RPS17    | 15q25.2  | 180472 | Ribosomal Protein S17                                                       | Diamond-Blackfan anemia 4                      |
| RPS17L1  | 15q25.2  | 180472 | Ribosomal protein S17a-Like 1                                               | Diamond-Blackfan anemia 4                      |
| AP3B2    | 15q25.2  | 602166 | Adaptor-Related Protein Complex 3, Beta-2 Subunit                            | Epileptic encephalopathy, early infantile, 48  |
| HOMER2   | 15q25.2  | 604799 | Homer, Drosophila, Homolog Of, 2                                            | Deafness, autosomal dominant 68                |
| WDR73    | 15q25.2  | 616144 | Wd Repeat-Containing Protein 73                                              | Galloway-Mowat syndrome 1                      |
| ALPK3    | 15q25.3  | 617608 | Alpha Kinase 3                                                              | Cardiomyopathy, familial hypertrophic 27       |
| SLC28A1  | 15q25.3  | 606207 | Solute carrier family 28, member 1                                          | Uridine-cytidineuria                           |
| AGBL1    | 15q25.3  | 615496 | Atp/Gtp-Binding Protein-Like 1                                              | Corneal dystrophy, Fuchs endothelial, 8         |
The clinical manifestations presented here were linked to 15q duplication in the fetus detected by CMA. As a molecular genetics detection technique, CMA can detect many chromosome structural abnormalities, but the technology has limitations. The combination and rational application of cytogenetics and molecular genetics technologies will undoubtedly open up the field of clinical application for such anomalies.

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CONFLICT OF INTEREST
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