Case report

t (3; 22)(q21;q12) in a pregnant woman with two abortions and postnatal death of one offspring

Zia Eslami¹ M.D., Mohammad Hasan Sheikhha² Ph.D., Seyed Mehdi Kalantar² Ph.D., Seyed Mohammad Seyedhasani² M.D.

- 1 Department of Pediatric, Yazd Shahid Sadoughi University of Medical Sciences, Yazd, Iran.
- 2 Department of Genetics, Research and Clinical Centre for Infertility, Yazd Shahid Sadoughi University of Medical Sciences, Yazd, Iran.

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Abstract

Background: Carriers of translocations may have an increased risk of an unbalanced progeny due to imbalances and delays in meiosis.

Case: A 24-year-old pregnant Iranian female was referred to the Genetic Department of Yazd Clinical and Research Centre for Infertility because of her pregnancy history. She had three previous pregnancies, two of which ended in abortion. The one live born infant was a girl who had multiple abnormalities and died when she was 11 days old. The cytogenetic analysis showed that the woman is a carrier of chromosomal translocation 46, XX, t (3; 22) (q21; q12), while her husband's karyotype was found to be normal. The karyotype of her mother showed the same translocation. The risk of further miscarriages was high, and the proband was monitored closely during her pregnancy. After nine months of pregnancy, a normal baby girl weighted 3460 gr was delivered by Caesarean section. Three hours after birth, the baby suffered from jaundice and respiratory distress. The baby's phenotype was normal. She received routine treatment successfully and after 15 days she was discharged from the hospital in a good condition. The baby's karyotype showed the same translocation as her mother and grandmother.

Conclusion: To our knowledge, no translocation with such breakpoints t (3; 22) (q21; q12) has been described previously in the women with RPL.

Key words: t(3;22), Karyotype, Miscarriage, Translocation.

Introduction

Recurrent spontaneous abortion, defined as three consecutive abortions, occurs in approximately 1% to 2% of couples. Although the cause is unknown in up to 50% of cases, about 5% of these couples are found to be a balanced translocation carrier(1). A translocation carrier may have an increased risk of unbalanced progeny. As a result, miscarriages frequently occur. In fact, many

Corresponding author:

Dr Mohammad Hasan Sheikhha, Genetics Department, Clinical and Research Center for Infertility, No.2, BouAli Avenue, Safayeh, Yazd, Iran. e-mail: sheikhha@yahoo.com carriers of translocations have a risk of miscarriages of about 5-10%. The risk will be higher in case of duplication/deletion of larger regions (2). In addition, chromosome aberrations are found in 27% of couples with fertility problems (3). In one study, Stephenson and Sierra (2006) identified 51 carriers of a structural chromosome rearrangement in 1893 couples (2.7%) with repeated pregnancy loss (RPL) (4).

Case Report

A pregnant woman was referred for genetic counselling, having had two first trimester abortions (after 6–8 weeks of pregnancy) and one

postnatal death of her offspring. This offspring was a girl who had multiple abnormalities including cleft lip and cleft palate, glaucoma and blindness and died when she was 11 days old. Chromosomal analyses had not been done on any of the aborted foetuses or died offspring which had all occurred prior to the cytogenetic analysis of the mother. The infertility of the couple was previously regarded as idiopathic, as none of the common reasons of infertility was found. The husband was a healthy, 30 year old, phenotypically normal male. His semen analysis showed a normal sperm count. His only sister had died for unknown causes when she was 6 months old. His mother and his father were cousin. The proband had four normal and healthy sisters and no brother. Her mother was not related to her father and she had only one sisters. Her mother had a cousin with the history of five repeated pregnancy losses and one normal daughter. There had been no other history of repeated spontaneous abortions in the family. Karoytyping was done for the couple. Chromosomal analyses were done on preparations made from peripheral blood by use of standard methodologies. G banding (5) was done and 20 metaphases spreads were analysed from the cytogenetic examination couple. The performed on cultured lymphocytes with a resolution of ~550. G banding of chromosomes derived from the woman's lymphocyte cultures showed a balanced translocation of chromosomes 3 and 22, breakage and reunion have occurred at bands 3q21 and 22q12 and the segments distal to these bands have been exchanged, thus the karyotype was 46,XX,t(3;22)(q21;q12). (Figure 1) Her husband was found to have a normal male karyotype.

Examination of peripheral blood lymphocytes obtained from the proband's mother showed the same translocation in the mother as well. Her father showed a normal karyotype 46, XY. The translocation was found in some other family members as well (Figure 2). The translocation was first interpreted by conventional cytogenetic analysis as a t(3;22)(q21;q12) inherited from the mother. The risk of further miscarriages was high, and the proband was monitored closely during her pregnancy. After nine months of pregnancy, a normal baby girl weighted 3460 gr was delivered by Caesarean section. Three hours after birth, the baby suffered from jaundice and respiratory distress. The hyperbilirubinemia was investigated and it appeared to be a result of ABO incompatibility. It was treated by exchange transfusion and phototherapy. In addition, a systolic murmur (grade II/VI) was heard on chest auscultation and the echocardiography revealed right ventricular cardiomyopathy, which appeared to be secondary to haemolysis and anaemia, as after correction of anaemia the murmur was disappeared and repeated echocardiography became normal. The baby was discharged from the hospital after 15 days in a good condition. The baby's karyotype showed the same translocation as her mother and grandmother.

The family pedigree was prepared according to Rimoin *et al.* (6). The known karyotypes of the family members were: the proband 46,XX,t(3;22); proband's husband 46,XY; proband's mother 46,XX,t(3;22); proband's father 46,XY; and the proband's daughter 46,XX,t(3;22), proband's aunt and cousin 46,XX,t(3;22), and cousin of proband's mother 46,XX,t(3;22). Other family members either were not cooperated or were not available to participate in cytogenetic examinations (Figure 2).

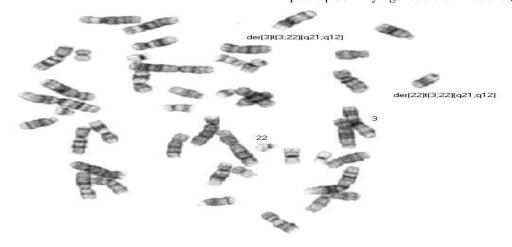


Figure.1: The karyotype of proband showing 46, XX, t (3;22)(q21;12).

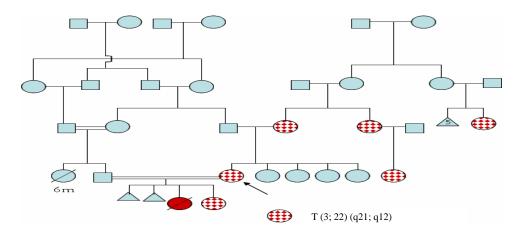


Figure 2: The family pedigree of the case

Discussion

Carriers of balanced aberrations have an increased risk of an unbalanced progeny due to imbalances and delays in meiosis (2). There are only few publications presenting translocations involving 3q and 22q. While, there are reports of other translocation in RPL. Joseph and Thomas (1987) reported a balanced reciprocal translocation between chromosomes 1 and 7 in both the husband and his mother in a couple who referred for chromosomal analysis having had three first trimester abortions (7). In addition, defects in chromosomes 3 and 22 are reported to be related to other pathological conditions. Partial trisomies of 22q were reported in connection with congenital abnormalities (8-10). Monosomy 22q was also described resulting in serious dysmorphic features and psychomotor delays (11-12). Translocation t(3;22)(q27;q11) was reported to be related with diffuse non-Hodgkin's lymphoma (13). In addition, T-prolymphocytic leukemia, small cell variant, associated with complex cytogenetic findings including t(3;22)(q21;11.2) (14).

To our knowledge, no translocation with such breakpoints t(3;22)(q21;q12) has been described previously in the women with RPL. The abnormal karyotype of the proband 46, XX, t (3; 22) (q21; q12) can be regarded as a reason of fertility problems in the investigated couple. The risk of further miscarriages is high due to large segments of duplication/deletion caused by unequal segregation. In the case that the couples seek more pregnancy, close monitoring is necessary. In addition more advanced techniques including prenatal diagnosis (PND) and preimplantation

genetic diagnosis (PGD) are advisable. These techniques for people suffering recurrent miscarriages are increasingly being performed worldwide (15). In the cases that PND is performed, if the fetus suffers the same balanced translocation as his/her mother, the pregnancy should be continued, but in the cases of larger duplication/deletion, it must be terminated.

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