Expectant management and live birth outcomes for male balanced-translocation carriers

Two case reports and a literature review

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Abstract

Rationale: Couples with male balanced-translocation carriers may experience recurrent pregnancy loss (RPL). Although the expectant management of RPL has developed over many years, genetic counseling for RPL couples with male balanced-translocation carriers remains challenging. Here, we describe the expectant management of 2 male carriers of balanced translocations.

Patient concerns: A 32-year-old and a 28-year-old man presented at the clinic with diagnoses of infertility following spontaneous abortions by their wives.

Diagnosis: Both patients had normal semen diagnosed by routine semen analysis and underwent cytogenetic diagnoses.

Interventions: Following genetic counseling and informed consent, both couples voluntarily chose expectant management with natural conception.

Outcomes: One couple experienced 2 natural pregnancies, the first of which ended in spontaneous abortion and the second produced a phenotypically normal infant. The other couple’s first pregnancy resulted in a fetus with a balanced translocation confirmed by amniocentesis and cytogenetic analysis.

Lessons: Expectant management with natural conception may be an alternative to genetic counseling in male balanced-translocation carriers with RPL, especially those who are reluctant to undergo preimplantation diagnosis.

Abbreviations: PGD = preimplantation genetic diagnosis, RPL = recurrent pregnancy loss.

Keywords: balanced translocation, expectant management, live birth, recurrent pregnancy loss

1. Introduction

Chromosomal translocation is an important cause of genetic changes in humans. Reciprocal translocations are the most common structural rearrangement in infertile men. Male translocation carriers often exhibit reproductive issues such as male infertility or recurrent pregnancy loss (RPL) in their spouses. RPL is an obstetric complication that affects male infertility or recurrent pregnancy loss (RPL) in their translocation carriers often exhibit reproductive issues such as male infertility or recurrent pregnancy loss. A 32-year-old and a 28-year-old man presented at the clinic with diagnoses of infertility following spontaneous abortions by their wives.

Large studies have suggested 2 options for male reciprocal-translocation carriers who experience RPL. First, preimplantation genetic diagnosis (PGD) is recommended as a tool to improve live birth rates and reduce the rate of miscarriage. Fischer et al  reported that translocation carriers who experienced 3 or more losses benefited from PGD, with an increased pregnancy-success rate, reduced length of time to conceive, and reduced pregnancy-loss rate. Sugiura-Ogasawara et al  reported that PGD could benefit translocation carriers by reducing the risk of miscarriage and avoiding a pregnancy with an unbalanced form of the translocation. In clinical practice, parental carriers of chromosomal translocations and a history of RPL are more likely to pursue a natural pregnancy following natural conception, and the cumulative live-birth rate with natural conception was reported to be 65% to 83%. Expectant management with natural conception has recently received attention for translocation carriers with RPL.

In clinical practice, parental carriers of chromosomal translocations and a history of RPL are more likely to pursue a natural pregnancy. Although PGD can reduce the abortion rate, it has a cost disadvantage. Furthermore, previous studies have shown no difference in reproductive outcomes, miscarriage rates, time to live birth, or live-birth rates between couples undergoing PGD and those pursuing a natural pregnancy. Hence, counseling for RPL patients with reciprocal translocation should include natural pregnancy or expectant management as treatment options.

This case report assesses the expectant management of 2 couples including men with chromosomal translocations, and further reviews its clinical application.
2. Case presentation

In these 2 cases, we carried out clinical expectant management for carriers of balanced translocations in couples with recurrent spontaneous abortions. This report was approved by the Ethics Committee of the Second Hospital, Jilin University, and written informed consent was obtained from both patients.

2.1. Case 1

A 32-year-old man visited the andrology service in March 2017 because his wife had experienced 2 spontaneous abortions before 13 weeks of gestation after 4 years of marriage. The patient had normal appearance and intelligence. Semen analysis revealed normal sperm concentration, motility, and morphology. The patient underwent cytogenetic detection, which revealed a karyotype of 46,XY,t(3;6)(q23;p21.3) (Fig. 1A). His wife’s karyotype was 46,XX. Following genetic counseling, the couple refused PGD because of family and financial conditions, and chose to pursue natural conception. We considered expectant management as a possible option, and the couple provided informed consent for expectant management treatment. To improve the couple’s confidence, we reassured them that balanced-translocation carriers can have natural pregnancies and produce phenotypically normal children. The couple received further attention before and during pregnancy, but the first pregnancy unfortunately resulted in spontaneous abortion at 10 weeks of gestation. However, a second pregnancy after 1 year passed 13 weeks of gestation successfully. Amniocentesis performed at 18 weeks of gestation showed that the fetus was a balanced-translocation carrier, consistent with the father’s karyotype. A phenotypically normal child was subsequently born.

2.2. Case 2

An apparently normal 28-year-old man presented in April 2018 with a 3-year history of primary infertility, after his wife had experienced 3 spontaneous abortions before 13 weeks of gestation after 4 years of marriage. The patient had normal appearance and intelligence. Semen analysis revealed normal semen quality. The cytogenetic results revealed his karyotype as 46,XY,t(6;11)(q21;q25) (Fig. 1B). His wife’s karyotype was 46, XX. Following genetic counseling, the couple provided informed consent for expectant management treatment. Their first attempted natural pregnancy passed 13 weeks safely, and amniocentesis and cytogenetic analysis at 17 weeks of gestation showed a fetus with a balanced translocation. This infant was subsequently delivered successfully.

2.3. Literature review

We searched for reports on expectant management of patients with chromosome translocations and RPL in PubMed using the keywords “expectant management/recurrent pregnancy loss.” Cases of chromosomal translocation were collected. A total of 11 studies involving live births following natural pregnancies were found. The reproductive outcomes of expectant management and natural conception in couples with recurrent abortion reported in previous studies are shown in Table 1. The live-birth rate for couples following natural conception was 25% to 71%.

3. Discussion

In this study, we report the expectant management outcomes in 2 cases of RPL in couples with male balanced-chromosomal translocation carriers. One male carrier had the reciprocal translocation 46,XY,t(3;6)(q23;p21.3) and the other was 46,XY,t(6;11)(q21;q25). Parental chromosome rearrangements are one of the main causes of RPL, and the rates of pregnancy loss are higher in carriers compared with those with RPL and normal karyotypes. Balanced reciprocal translocations are the most common structural rearrangement. Men affected by such translocations may have failure of spermatogenesis and/or RPL. It has been reported that specific chromosomes and breakpoints involved in translocation are related to RPL and the involved chromosomes and breakpoints should be considered in genetic counseling.

However, pregnancy resulting in live birth is possible in translocation carriers with RPL without treatment. Page et al. reported that the rate of live births in carriers with balanced translocations was much higher than anticipated, with a live-birth rate of about 60% to 70% without treatment, depending on the specific chromosome or breakpoint. In addition, the pregnancy, live-birth, and clinical miscarriage rates were similar in patients treated with expectant management and those treated with PGD. Expectant management also has the benefit of lower cost per live birth, and expectant management is thus a worthy clinical option for balanced-translocation carriers.

Despite developments in expectant management, there are still challenges in relation to translocation carriers with RPL. This study explored the expectant management of 2 male carriers with RPL, involving chromosomes 3, 6, and 11 and breakpoints 3p23, 6p21.3, 6q21, and 11q25. Live births have previously been reported for male carriers with chromosome t(3;6)(q12;q27). In addition, familial balanced translocation carriers with t(3;6)(q12; q15) or t(3;6)(p12.3;q24.3) have frequently been reported, suggesting that their reproductive function is normal. Natural conception may be a more viable option for fertile carriers of translocations with a low risk of conceiving a chromosomally unbalanced offspring. To the best of our knowledge, this is the first report of the expectant management of carriers of chromosomal translocations with RPL in the Chinese population.
| No | PDG | Reproductive outcomes of expectant management or natural conception for the couples with recurrent abortion reported in previous literature. | Reference |
|----|-----|----------------------------------------------------------------------------------------------------------------------------------|------------|
| 1  |     | Median time to pregnancy was 6.5 mo. The pregnancy rate, live birth rate and clinical miscarriage rate were similar with PGS group. | Murugappan et al[92] |
| 2  |     | The live-birth rate was 53%, and the cost was $45,300 per live birth. IVF/PGS was not a cost-effective strategy for increasing live birth. | Murugappan et al[93] |
| 3  | NA  | Mean time to live birth was 23.3 months in PDG. The live birth rate was 66.6%. | Mahthirpalai et al[94] |
| 4  | NA  | Following treatment, 35.6% of the couples had a healthy live-born child. | Flynn et al[95] |
| 5  | NA  | The live birth rates on the first PDG trial were 37.8%. | Ikuma et al[96] |
| 6  | NA  | The live birth rates on the first natural pregnancy 53.8%. | Sugiura-Ogasawara et al[97] |
| 7  | NA  | Patients with parental chromosomal rearrangements do not have a significantly lower live birth rate than patients without aberrations. | Carpi et al[98] |
| 8  | NA  | The live birth rate was 25% to 71% among 847 couples who conceived naturally. | Kong et al[99] |
| 9  | NA  | The live birth rate was 71% among 58 monitored pregnancies. | Stephenson et al[100] |

NA = not available, PDG = preimplantation genetic diagnosis.

4. Conclusions

In conclusion, this study reported 2 couples including male balanced-translocation carriers who received expectant management. Natural conception with expectant management may be alternatives to genetic counseling in these patients, especially those who are reluctant to undergo PGD.

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References

[1] Mayeur A, Abdad N, Hesters L, et al. Chromosomal translocations and semen quality: a study on 144 male translocation carriers. Reprod Biomed Online 2019;38:46–55.
[2] Zhang HG, Wang RX, Pan Y, et al. A report of nine cases and review of the literature of infertile men carrying balanced translocations involving chromosome 5. Mol CytoGenet 2018;11:10.
[3] Sundheimer LW, Liu L, Baysal RP, et al. Diagnosis of parental balanced reciprocal translocations by trophectoderm biopsy and comprehensive chromosomal screening. J Assist Reprod Genet 2018;35:163–9.
[4] Priya PK, Mishra VV, Roy P, et al. A study on balanced chromosomal translocations in couples with recurrent pregnancy loss. J Hum Reprod Sci 2018;11:337–42.
[5] Kohn TP, Kohn JR, Darilek S, et al. Genetic counseling for men with recurrent pregnancy loss or recurrent implantation failure due to abnormal sperm chromosome aneuploidy. J Assist Reprod Genet 2016;33:571–6.
[6] Zhang X, Zhang H, Hu C, et al. Clinical features of carriers of reciprocal chromosomal translocations involving chromosome 2: report of nine cases and review of the literature. Int Braz J Urol 2018;44:785–93.
[7] Iews M, Tan J, Taskin O, et al. Does preimplantation genetic diagnosis improve reproductive outcome in couples with recurrent pregnancy loss owing to structural chromosomal rearrangement? A systematic review. Reprod Biomed Online 2018;36:677–85.
[8] Fischer J, Collis P, Escudero T, et al. Preimplantation genetic diagnosis (PGD) improves pregnancy outcome for translocation carriers with a history of recurrent losses. Fertil Steril 2010;94:283–9.
[9] Scriven PN, Flinter FA, Khalaf Y, et al. Benefits and drawbacks of preimplantation genetic diagnosis (PGD) for reciprocal translocations: lessons from a prospective cohort study. Eur J Hum Genet 2013;21:1035–41.
[10] Flynn H, Yan J, Saravelos SH, et al. Comparison of reproductive outcome, including the pattern of loss, between couples with chromosomal abnormalities and those with unexplained repeated miscarriages. J Obstet Gynaecol Res 2014;40:109–16.
[11] Sugiuра-Ogasawara M, Aoki K, Fujii T, et al. Subsequent pregnancy outcomes in recurrent miscarriage patients with a paternal or maternal carrier of a structural chromosome rearrangement. J Hum Genet 2008;53:622–8.
[12] Ikuma S, Sato T, Sugiuра-Ogasawara M, et al. Preimplantation genetic diagnosis and natural conception: a comparison of live birth rates in patients with recurrent pregnancy loss associated with translocation. PLoS One 2015;10:e0129958.
[13] Mahthirpalai S, Durland U, Havelock J, et al. Prevalence and treatment choices for couples with recurrent pregnancy loss due to structural chromosomal anomalies. J Obstet Gynaecol Can 2018;40:653–62.
[14] De Krom G, Arens YH, Coonen E, et al. Recurrent miscarriage in translocation carriers: no differences in clinical characteristics between couples who accept and couples who decline PGS. Hum Reprod 2015;30:484–9.
[15] Franssens MT, Musters AM, van der Veen F, et al. Reproductive outcome after PGD in couples with recurrent miscarriage carrying a structural chromosome abnormality: a systematic review. Hum Reprod Update 2011;17:467–75.
[16] Page JM, Silver RM. Genetic causes of recurrent pregnancy loss. Clin Obstet Gynecol 2016;59:498–508.
[17] Zhang H, Wang R, Li L, et al. Clinical feature of infertile men carrying balanced translocations involving chromosome 10: case series and a review of the literature. Medicine (Balitmore) 2018;97:e0452.
[18] Pal AK, Ambulkar PS, Waghmare JE, et al. Chromosomal aberrations in couples with pregnancy loss: a retrospective study. J Hum Reprod Sci 2018;11:247–53.

[19] Zhang HG, Liu XY, Hou Y, et al. Reproductive outcome of a case with familial balanced translocation t(3;6): implications for genetic counseling. Genet Mol Res 2015;14:2809–15.

[20] Yusenko MV, Nagy A, Kovacs G. Molecular analysis of germline t(3;6) and t(3;12) associated with conventional renal cell carcinomas indicates their rate-limiting role and supports the three-hit model of carcinogenesis. Cancer Genet Cytogenet 2010;201:15–23.

[21] Eleveld MJ, Bodmer D, Merks G, et al. Molecular analysis of a familial case of renal cell cancer and a t(3;6)(q12;q15). Genes Chromosomes Cancer 2001;31:23–32.

[22] Murugappan G, Shahine LK, Perfetto CO, et al. Intent to treat analysis of in vitro fertilization and preimplantation genetic screening versus expectant management in patients with recurrent pregnancy loss. Hum Reprod 2016;31:1668–74.

[23] Murugappan G, Ohno MS, Lathi RB. Cost-effectiveness analysis of preimplantation genetic screening and in vitro fertilization versus expectant management in patients with unexplained recurrent pregnancy loss. Fertil Steril 2015;103:1215–20.

[24] Carp H, Feldman B, Oelsner G, et al. Parental karyotype and subsequent live births in recurrent miscarriage. Fertil Steril 2004;81:1296–301.

[25] Kong GW, Lok IH, Yiu AK, et al. Clinical and psychological impact after surgical, medical or expectant management of first-trimester miscarriage—a randomised controlled trial. Aust N Z J Obstet Gynaecol 2013;53:170–7.

[26] Stephenson MD, Sierra S. Reproductive outcomes in recurrent pregnancy loss associated with a parental carrier of a structural chromosome rearrangement. Hum Reprod 2006;21:1076–82.