Pre-pregnancy cytogenetic analysis of general couples in eastern China

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The aim of this study was to investigate the contribution of chromosomal anomalies and the frequency of particular types of aberrations in general couples preparing for pregnancy and make recommendations for pregnancy on the basis of the medical literature. A total of 6,198 general couples were included in the present study. The karyotypes were generated from the peripheral blood lymphocyte cultures and the cytogenetic analysis was performed using G-banding. In 12,396 cases, chromosomal anomalies were detected in 59 cases (0.48%, 59/12,396). Among of them, the frequency of translocation was 0.35% (n = 43). Sex chromosomal anomalies accounted for 0.07% (n = 9), including Klinefelter syndrome (KS) (n = 4), Turner syndrome (TS) (n = 4), and XYY syndrome (n = 1). The others, including inversions (n = 6) and deletion (n = 1), accounted for 0.06%. Our study indicates that clinically important chromosomal defects are present at a remarkable frequency in the general couples in eastern China, suggesting pre-pregnancy cytogenetic analysis should be routinely performed among general couples in this area so that informed decision can be made, which will help to improve the quality of the pregnancy.

Generally, the couples planning for their first pregnancy have no information related to reproductive problems. The reproducing disorders are the relatively common problem that affects couples worldwide. It is estimated that approximately one in six couples experiences difficulties in reproducing¹. It was reported that the frequency of chromosomal aberrations is approximately 8% in cases suffering reproductive failure such as infertility and pregnancy losses², while it is 0.6% in general population³, the majority of whom wouldn’t do cytogenetic analysis before encountering reproductive problems. In Zhenjiang City in eastern China, the cradle of the Chinese Wu culture and the region of the fastest-growing economy in China, the local government sponsors every couple for cytogenetic karyotype analysis before pregnancy if they would like to. The couples can chose one of the following three government-assigned institutes to do cytogenetic analysis: Zhenjiang Family Planning Guidance Institute, Zhenjiang Jingkou District Family Planning Guidance Station, and Department of Reproductive & Genetics, the Fourth Affiliated Hospital of Jiangsu University. Here, we report the contribution of chromosomal anomalies and the frequency of particular types of aberrations in general couples preparing for pregnancy, some suggestions are also made on the basis of the present study combining the medical literatures.

Results
A total of 6,198 couples (i.e. 12,396 cases) were evaluated during 1990–2010. Chromosome anomalies were found in 59 cases (0.48%, 59/12,396) (Table 1). Sex chromosome disorders were found in nine cases (15.3%, 9/59), including eight aneuploidy cases and one isochromosome of Xq (Table 2). Translocations were detected in 43 cases (72.9%, 43/59), including 33 balanced translocations and 10 robertsonian translocations. The other structural chromosomal anomalies, including inversions and deletion, were seen in seven cases (11.9%, 7/59) (Table 3). Totally, the incidence of the sex chromosome disorders and the structural chromosomal abnormalities were 0.08% (9/12,396) and 0.4% (50/12,396), respectively.

Discussion
The present study investigated the incidence of chromosomal abnormalities in 6,198 couples i.e. 12,396 cases. Overall, 0.48% of the cases were found to have chromosomal aberration. Chromosomal disorder is closely related to reproductive abnormalities according to previous literatures²–⁶. Chromosomal abnormalities, such as trisomy, reciprocal translocation, Robertsonian translocation, chromosomal inversion, and chromosomal deletion, could be the main factors related to the bad reproductive ending event². The goal of pre-pregnancy cytogenetic tests is to
uncover potential genetic defects in the prospective parents, thus helping them make informed decision about the planned pregnancy. For example, balanced reciprocal translocations carriers should undergo prenatal diagnosis to rule out any monosomies and trisomies in their offspring.

Klinefelter syndrome (KS) is one of the common causes of human male infertility. This study detected four cases of KS in 6,198 couples and the rate is about one in 1,500 (4/6,198) in the male. The patients with KS carry an additional X chromosome, which results in male hypogonadism, androgen deficiency, and impaired spermatogenesis. Treatment consists of testosterone replacement therapy to correct the androgen deficiency and to provide patients with appropriate virilization. This therapy also has positive effects on mood and self-esteem and has been shown to protect against osteoporosis, although it will not reverse infertility. Nowadays, however, patients with KS, including the non-mosaic type, should no longer be considered irreversibly infertile, because a biological pregnancy can be obtained by Testicular Sperm Extraction and Intra-Cytoplasmatic sperm injection (TESE-ICSI). TESE will be followed by cryopreservation of extracted sperms before introducing androgen treatment in adolescent and will be used latter for ICSI when the KS patient would like to seek children. But in some cases, chromosomal errors might be transmitted to the offspring of males with this syndrome. The genetic implications of the fertilisation procedures, including pre-transfer or prenatal genetic assessment, should be explained to patients and their spouse.

In the current study, TS was detected in four cases, one of which is a isochromosome 46,X, i(Xq), with the rate of one in 1500 (4/6,198) in the female that is higher than one in 3000 to one in 2500 in live-born girls reported in the literature. We only dealt with the cases of TS the female that is higher than one in 3000 to one in 2500 in live-born

| Abnormalities | Cases[n] |
|---------------|----------|
| Aneuploids    | 8        |
| Isochromosome of Xq | 1        |
| Translocations | 43       |
| Inversions    | 6        |
| Deletion      | 1        |
| **Total**     | 59       |

Table 1 | Total chromosome abnormalities among 6,198 general couples

Most XYY syndromes have baby, but the same chromosome errors might occur in the offspring, which suggested that the prenatal diagnosis should be performed after pregnancy.

Reciprocal translocations, typically an exchange of two terminal segments from different chromosomes, will lead to recurrent miscarriages leading to bad obstetric history; for others, there may be a risk of offspring with mental and physical disability due to segregation of the translocation chromosomes at meiosis resulting in sperm or eggs with chromosome imbalance.

Conventional karyotyping can be used for analyzing fetal chromosomes including distinguishing the balanced carriers or the fetus with unbalanced translocations through amniocentesis, chorion villus sampling (CVS) or periumbilical cord blood sampling (PUBS) after their natural pregnancy. The advents of in-vitro fertilization (IVF) or intracytoplasmic sperm injection (ICSI) have enabled people to increase the pregnancy chance of the chromosome balanced translocations carriers. Preimplantation genetic diagnosis (PGD) has been very successful in assisting couples to achieve a viable unaffected pregnancy and has certainly reduced the time taken to achieve a pregnancy. PGD is used for reciprocal translocation carriers to minimize the risk of having an affected child or the distress of pregnancy termination, and to reduce the risk of miscarriage due to abnormal segregation of the translocation.

Robertsonian translocation is a special kind of balanced translocation, involving only the acrocentric chromosomes, specifically chromosomes 13, 14, 15, 21 and 22. They are formed when the long arms of two acrocentric chromosomes are translocated, ultimately forming a single chromosome. The clinical significance of the robertsonian translocation is the increased risk of infertility, spontaneous abortion and unbalanced chromosomal offspring. In the current study, we found three cases of robertsonian translocation involving chromosome 21, accounting for 30% (3/10) of the sex chromosome disorders. The potential live-born unbalanced outcome of this robertsonian translocation is translocation trisomy 21, resulting in Down syndrome.

Reproductive risks of Robertsonian translocations are dependent on the chromosomes involved. For couples with robertsonian translocations, the prenatal diagnosis should be suggested after their natural pregnancy to avoid the serious fetal malformation like trisomies. Like reciprocal translocation, robertsonian translocation couples need assisted conception for subfertility. Therefore, PGD should be performed after careful risk assessment and genetic counselling.

The government policy that recommends and offers free pre-pregnancy cytogenetic diagnosis to all willing couples has enabled us to uncover diverse and rather frequent chromosomal abnormalities “hidden” in the couples planning for pregnancy. Obviously, this policy not only achieved a remarkable success but also raised many interesting questions: How many of these chromosomal syndromes would be probably detected by this way? How many people would benefit from the management after the sponsored cytogenetic analysis? How about the cost/benefit of this policy? We will attempt to address these questions in our future studies.

Methods

Study population. A retrospective study was conducted in the couples planning for pregnancy during the period of 1990 to 2010 in Zhenjiang City, eastern China. We carefully screened our study subjects, ensuring that the couples enlisted had no prior examination or heath history, suggestive of fertility problems. A total of 6,198 couples, namely 12,396 individuals, were included. The average age of females is 23 ± 4, and the males is 26 ± 3; in our region, the average maternal age in last decade is about 26 ± 6, according to the government statistics calculating by Zhenjiang Family Planning Guidance institute. Informed consent was obtained from all subjects in the present study.

Chromosomal analysis. Karyotypes were generated from the peripheral blood lymphocyte cultures, and the metaphase chromosome preparations were made according to standard cytogenetic protocols. Cytogenetic analysis was performed by
### Table 3 | Structural chromosomal abnormalities identified in this study

| Structural anomalies | Males | Females |
|----------------------|-------|---------|
| Reciprocal translocations |       |         |
| 46, XX, (4;10)(q35;q24) | 1     |         |
| 46, XY, t(1;2)(p13;p13) | 1     |         |
| 46, XX, rob(14;21)(q10;q10) | 2     |         |
| 46, XY, rob(14;14)(q10;q10) | 1     |         |
| 46, XX, inv(10)(p11.2;p23) | 1     |         |
| 46, XY, inv(3)(p14;q21) | 1     |         |
| 46, XX, t(5;14)(q13;q32) | 1     |         |
| 46, XY, t(Y;2)(q21;p23) | 1     |         |
| 46, XX, rob(13;14)(q10;q10) | 1     |         |
| 46, XY, rob(13;13)(q10;q10) | 3     |         |
| 46, XX, inv(1)(p36;q21) | 1     |         |
| 46, XY, inv(10)(p11.2;p14) | 1     |         |
| 46, XX, t(11;14)(q24;q12) | 1     |         |
| 46, XX, t(1;6)(q44;q13) | 1     |         |
| 46, XX, rob(15;22)(q10;q10) | 2     |         |
| 46, XX, inv(7)(p13;q22) | 1     |         |
| 46, XY, inv(2)(p22;q31) | 1     |         |
| 46, XX, t(1;2)(q31;p21) | 1     |         |
| 46, XY, t(1;9)(q23;q22) | 1     |         |
| 46, XX, t(3;11)(q21;q23) | 1     |         |
| 46, XY, t(4;5)(q35;p15.1) | 1     |         |
| 46, XX, t(6;10)(q14;p12) | 1     |         |
| 46, XX, t(5;9)(q31;q22) | 1     |         |
| 46, XX, t(5;18)(q22;q22) | 1     |         |
| 46, XX, t(12;14)(q22;q32) | 1     |         |
| 46, XY, t(7;15)(q22;q24) | 1     |         |
| 46, XX, t(2;9)(q21;p22) | 1     |         |
| 46, XX, t(3;5)(p21.3;q31) | 1     |         |
| 46, XX, t(9;10)(q22;q22) | 1     |         |
| 46, XY, t(4;9)(p14;q22) | 1     |         |
| 46, XX, t(2;9)(p12;q22) | 1     |         |
| 46, XY, t(2;3)(q13;q25) | 1     |         |
| 46, XX, t(4;17)(q31;q21) | 1     |         |
| 46, XY, t(7;20)(p22;p12) | 1     |         |
| 46, XX, t(13;14)(q32;q11.2) | 1     |         |
| 46, XX, t(3;14)(q12;q13) | 1     |         |
| 46, XX, t(14;18)(q23;q21) | 1     |         |
| 46, XX, t(8;21)(q24;q22) | 1     |         |

### Acknowledgments

The authors thank Zhenjiang Family Planning Guidance Institute and Jingkou District family planning guidance Station for the assistance with samples collecting. This work was supported by Jiangsu Research Project for the Key Talent of Maternal and Child Health under Grant No. FRC201124.

### Author contributions

Y.Y., H.W. and Y.T. designed, carried out the experiments, analyzed the data and prepared the draft of manuscript. M.G. and S.X. participated the experiments. X.X. and X.C. conceived the idea, supervised all research and revised the manuscript. All authors reviewed the manuscript.
Additional information

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Yang, Y. et al. Pre-pregnancy cytogenetic analysis of general couples in eastern China. Sci. Rep. 4, 7224; DOI:10.1038/srep07224 (2014).

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