Basic Human Genetics: Reproductive Health and Chromosome Abnormalities

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Training Course in Sexual and Reproductive Health Research
Geneva 2014
Categories of Genetic Diseases

- Single gene abnormalities
  - Autosomal dominant
  - Autosomal recessive
  - X-linked
  - Mitochondrial
  - Imprinting
- Chromosomal abnormalities
  - Numerical
  - Structural
  - Microdeletions
- Multifactorial diseases e.g. diabetes
- Acquired somatic abnormalities e.g. cancer
Types of Chromosome Abnormalities

Numerical:
- Trisomy
- Monosomy
- Mosaicism
- Triploidy

Structural:
- Translocation
- Deletion
- Inversion

Microdeletions and microinsertions
46 Chromosomes in a human cell as seen under the microscope
Frequencies of chromosome abnormalities

A chromosome abnormality is present in 40-50% of all recognized first-trimester pregnancy loss. Approximately 1 in 6 of all pregnancies results in spontaneous miscarriage.

Birth prevalence of chromosome abnormalities is 0.5-1%.
Chromosome abnormalities can cause:

- Infertility
- Repeated spontaneous abortions
- Stillbirths
- Infant mortality
- Birth defects
- Sexual ambiguity or abnormality in sexual development
- Unexplained short stature in female children
- Intellectual disability
Trisomy

Presence of an extra chromosome, the total number of chromosomes is 47 in a somatic cell. Trisomy usually results from meiotic non-disjunction.

There are 3 of number 21 chromosomes
Monosomy

- Absence of one chromosome, so the total number of chromosomes is 45 chromosomes in a somatic cell.
- Usually only seen as 45,X, (autosomal monosomy is usually lethal).
- Monosomy usually results from meiotic non-disjunction.
Mosaicism

There are 2 types of cells in an individual, for example normal 46,XY cells and abnormal trisomic cell line 47,XY,+21. The two cell lines are derived from the same zygote due to mitotic non-disjunction.
Triploidy

Presence of 3 haploid sets: $23 \times 3 = 69$ chromosomes
(haploid set = 23, diploid set = 46).

Usually incompatible with life and seen only in abortions.
May result from 2 sperms fertilising the ovum or retainment of the polar body with the ovum.
Normal meiosis, the 46 chromosomes become 23 in each gamete
Non-disjunction during meiosis means that one daughter cell gets 24 chromosomes and the other 22 chromosomes.
Fertilisation of the gamete carrying 24 chromosomes (extra number 21) with a normal gamete having 23 chromosomes results in a zygote of 47 chromosomes (trisomy 21 zygote).
Reciprocal translocation = exchange of segments between 2 non-homologous chromosomes
Robertsonian translocation occurs between 2 acrocentric chromosomes with breaks near centromeres and union of the long arms.
Carriers of balanced translocations are healthy but...

They are at risk of having offspring with unbalanced chromosome constitution

This may present as:
- Repeated spontaneous abortions
- Stillbirths
- Birth defects
- Intellectual disability
Deletion: loss of part of a chromosome
Pericentric inversion: two breaks with inversion of the segment in between
Karyotype description

- 46,XX normal female karyotype
- 46,XY normal male karyotype
- 45,X monosomy X = Turner syndrome
- 47,XY,+21 trisomy 21 = Down syndrome
- 46,XY, 5p- deletion of part of short arm of chromosome 5 = Cri du Chat syndrome
- 46, XX, t(2;4)(q22;q23) translocation between long arms of chromosomes 2 and 4 with breakpoints at region 2 band 2 for chromosome 2 and region 2 band 3 for chromosome 4
Consequences of chromosome abnormalities

- Repeated spontaneous abortions: healthy carriers of translocations and inversions.
- Stillbirths and infant deaths: where the chromosome abnormality is very severe for example trisomy 13 and trisomy 18.
- Congenital disorders: for example Down syndrome, microdeletion syndromes.
Syndromes with chromosome abnormalities
# Chromosome abnormalities at birth

<table>
<thead>
<tr>
<th>Abnormality</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>Trisomy 13</td>
<td>0.2/1000 births</td>
</tr>
<tr>
<td>Trisomy 18</td>
<td>0.3/1000 births</td>
</tr>
<tr>
<td>Trisomy 21</td>
<td>1.5/1000 births</td>
</tr>
<tr>
<td>45,X</td>
<td>0.2/1000 female births</td>
</tr>
<tr>
<td>47XXX</td>
<td>1/1000 female births</td>
</tr>
<tr>
<td>47,XXY</td>
<td>1/1000 male births</td>
</tr>
<tr>
<td>47,XYY</td>
<td>1/1000 male births</td>
</tr>
<tr>
<td>Other unbalanced rearrangements</td>
<td>1/1000 births</td>
</tr>
<tr>
<td>Balanced rearrangements Total</td>
<td>3/1000 births</td>
</tr>
</tbody>
</table>
Down syndrome (DS)

The overall prevalence at birth is approximately 1 in 650 to 1 in 700 births.
May be higher in some countries where women continue to bear children at an advanced age.
Clinical features of DS

The most common finding in the newborn period is severe hypotonia.

Single palmar creases are found in 50% of Down syndrome children in contrast to 2-3% of the general population.

Congenital cardiac abnormalities are present in 40-45% of babies with Down syndrome.

Hypothyroidism.
Facial features of DS

• Upward sloping palpebral fissures
• Brushfield spots and bilateral epicanthic folds
• Small ears
• Protruding tongue
Affected children show a broad range of intellectual disability with IQ scores ranging from 25 to 75. The average IQ of young adults with Down syndrome is around 40 to 45.

Social skills are relatively well advanced and most children with Down syndrome are happy and very affectionate.
Adult height is usually around 150cm. In the absence of a severe cardiac anomaly, which leads to early death in 15-20% of cases, average life expectancy is 50-60 years. Most affected adults develop Alzheimer disease in later life due to dosage effect of the amyloid precursor protein gene.
Chromosome abnormalities in Down syndrome

95% of cases are trisomy 21, 47,XX,+21 (47,XY,+21), risk of having trisomy 21 increases with advanced maternal age.

4% are due to translocation between chromosome 21 and another acrocentric with a total number of chromosomes =46, but the genetic material of chromosome 21 is present in triplicate. The translocated chromosome is usually inherited from a normal carrier parent. Such a translocation carrier parent has a risk of having a Down syndrome with each pregnancy (about 20% if mother is carrier and 5% if father is carrier).

1% mosaic cases (46,XY/47,XY,+21).
<table>
<thead>
<tr>
<th>Maternal Age at Delivery (yr)</th>
<th>Risk of Down’s Syndrome</th>
<th>Risk of Any Chromosomal Abnormality</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>1/1667</td>
<td>1/526</td>
</tr>
<tr>
<td>25</td>
<td>1/1200</td>
<td>1/476</td>
</tr>
<tr>
<td>30</td>
<td>1/952</td>
<td>1/385</td>
</tr>
<tr>
<td>35</td>
<td>1/378</td>
<td>1/192</td>
</tr>
<tr>
<td>40</td>
<td>1/106</td>
<td>1/66</td>
</tr>
<tr>
<td>45</td>
<td>1/30</td>
<td>1/21</td>
</tr>
</tbody>
</table>

Salihu, 2003, Obstet Gynecol
Meiosis in Robertsonian translocation carrier who has 45 chromosomes (only one 21) but the other chromosome 21 is translocated to chromosome 22 = normal amount of genetic material
The gamete carrying 23 chromosomes but one is a translocation 21/22 is fertilised by a normal gamete resulting in a zygote with translocation Down syndrome.
Turner syndrome: monosomy X

The two main medical problems are short stature and ovarian failure. Ovarian failure leads to primary amenorrhea and infertility. Estrogen replacement therapy should be initiated at adolescence for the development of secondary sexual characteristics and long-term prevention of osteoporosis.
Normal female karyotype and Monosomy X
Turner syndrome features

- Lymphedema at birth
- Low posterior hair-line
- Increased carrying angles at the elbows
- Short fourth metacarpals
- Widely spaced nipples
- Coarctation of the aorta present in 15% of cases
Chromosome anomalies in Turner syndrome

45,X = 50%

Mosaics: 46,XX/45,X = 35%

Structural abnormalities (deletion, isochromosome, ring X)
Turner syndrome is being detected early in pregnancy as a result of routine detailed ultrasound scanning, which can reveal either generalized edema (hydrops) or swelling localized to the neck (nuchal cyst or thickened nuchal pad).
Klinefelter syndrome: 47,XXY

- Infertility.
- Hypogonadism.
- Diminished secondary sexual characters.
- Clumsiness or mild learning difficulties. The overall verbal IQ is reduced by 10-20 points below that of unaffected siblings and controls.
- 30% of adult males will show gynecomastia (enlargement of the breasts).
Chromosome abnormalities in Klinefelter syndrome

47,XXY
48,XXXXY, 48,XXYY
49,XXXXY
46,XY/47,XXY
XYY male

47, XYY

[Diagram of human chromosomes showing 47 chromosomes, including an extra Y chromosome (YY).]
XYY MALES

Fertility is normal.
Physical appearance is normal and stature is usually above average.
Intelligence is mildly impaired, with an overall IQ score of 10-20 points below a control sample.
The additional Y chromosome must arise as a result of non-disjunction in paternal meiosis II or as a post-zygotic event.
XXX females

These women usually have no physical abnormalities but can show a mild reduction of between 10 and 20 points in intellectual skills below their siblings. This is rarely of sufficient severity to require special education.

Women with a 47,XXX karyotype usually show normal fertility and have children with normal karyotypes.
Chromosome microdeletion syndromes
Prader-Willi syndrome

- Hypotonia
- Poor sucking and feeding in neonates
- Fair skin and hair
- Downturned mouth corners
- Hyperthermia
- Gestational history of diminished fetal movements
- Hyperphagia and obesity
- Short stature
- Small hands and feet
- Mental subnormality
- Narrow bifrontal diameter
- Hypogonadotropic hypogonadism
- Caused by microdeletion of paternal 15q11.2-12 (75%)
Diagnosis of microdeletion syndromes

FISH techniques using specific probes.
Array comparative genomic hybridisation (array CGH).
Microdeletions usually cannot be detected in the standard banded karyotype.
Prader-Willi Syndrome

Fluorescent in situ hybridization (FISH) demonstrating deletion (del) of SNRPN probe on one of the chromosomes 15s.

Figure 1
Indications for chromosome analysis in lymphocytes

- Multiple congenital abnormalities
- Unexplained mental retardation
- Sexual ambiguity or abnormality in sexual development
- Infertility
- Recurrent miscarriage
- Unexplained stillbirth
- Unexplained short stature in female children
- Malignancy and chromosome breakage syndromes
Conclusions

Around 20,000 chromosome abnormalities have been registered in laboratory databases.

Chromosome abnormalities contribute to about 8% of all birth defects.

Chromosome abnormalities can be diagnosed in the fetus through chorion villus biopsy or amniocentesis (prenatal diagnosis), or by preimplantation genetic diagnosis following IVF.