**Case Reports**


Trisomy of the short arm of chromosome 10

**Summary.** A case of a fetus with multiple malformations is described. The mother showed a 46,XX,rcp(10;22) (p11;p11) karyotype. Amniocentesis at the 16th week of gestation revealed that the male fetus had a der(22) chromosome—that is, he was trisomic for a large part of 10p (10pter→10p11).

Clinical findings of cases with 10p, 10q, and mosaic 10 trisomies are briefly reviewed.

Recently, Nakagome et al (1973) reported a case of mosaic 10 trisomy. In addition, they found two cases of C trisomy with strikingly similar clinical features reported in the literature. They postulated that the trisomy 10 was a clinical as well as cytogenetic entity. In the present report a case is described of a prenatally diagnosed 10p trisomy due to maternal rcp(10;22)(p11;p11) translocation. Necropsy findings are also presented.

**Case report**

A 30-year-old woman was referred to us for prenatal chromosome analysis. She had previously had a malformed boy and chromosome studies revealed that she, one of her two sisters, and her mother were carriers of a t(Cp−;Gp+) translocation (Yanagisawa and Adachi, 1970). The boy had a 46,XY,−G,+F karyotype—that is he was trisomic for the short arm of a C-group chromosome.

Since then the woman has had two pregnancies; however, both of them were terminated because of her fear of having another malformed baby. Recently, she learned about amniocentesis and elected to have the test.

Amniocentesis in the 16th week of gestation revealed that the fetus had the same unbalanced karyotype as the malformed brother. In the 22nd week of gestation, the pregnancy was terminated. The result of amniotic-

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![Fig. 1. Appearance of the fetus showing hypertelorism, cleft lip, and micrognathia.](http://jmg.bmj.com/ on June 20, 2017 - Published by group.bmj.com)
TABLE
SUMMARY OF CLINICAL FINDINGS IN CASES WITH 10, 10p, AND 10q TRISOMY

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<tbody>
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<td>Karyotype</td>
<td>+ 10/normal</td>
<td>+ 10p</td>
<td>+ 10p*</td>
<td>+ 10p†</td>
<td>+ 10q</td>
</tr>
<tr>
<td>Age</td>
<td>5 yr</td>
<td>Fet us (22 wk)</td>
<td>Died at 101 days of age</td>
<td>10 yr</td>
<td>Died at 2 days of age</td>
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<tr>
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<td>M</td>
<td>F</td>
<td>M</td>
<td>M</td>
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<tr>
<td>Mental retardation</td>
<td>+</td>
<td>?</td>
<td>?</td>
<td>+</td>
<td>?</td>
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<tr>
<td>Deformed skull</td>
<td>+</td>
<td>-</td>
<td>-</td>
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<tr>
<td>Strabismus</td>
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<td>?</td>
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<tr>
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<tr>
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<td>- ?</td>
<td>+</td>
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<tr>
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<td>+</td>
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<tr>
<td>Cleft palate</td>
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<td>+</td>
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<tr>
<td>Micrognathia</td>
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<td>+</td>
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<td>Agenesis of corpus callosum</td>
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<td>?</td>
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<tr>
<td>Arthrogryposis or flexion deformity of fingers and toes</td>
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<td>-</td>
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<tr>
<td>Deformed feet</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Abnormal genitalia</td>
<td>-</td>
<td>+</td>
<td>+</td>
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</tbody>
</table>

* Originally reported as partial C trisomy (see text).
† Originally reported as trisomy 12q (see text).

tertiary trisomy for a large part of the short arm of a No. 10 chromosome (10pter→10p11). A very small part of the short arm and the satellite of a No. 22 were missing.

Discussion
As far as the authors are aware, there has been only one reported case of trisomy 10 in which the extra chromosome was identified by banding studies (Nakagome et al, 1973). In 1967 Bühler et al described a case of ‘partial trisomy 12’. Recently, Tsuchimoto and Bühler (1974, personal communication) re-examined the case using banding techniques and found that the case was, in fact, trisomic for the distal part of 10q. They also described an unrelated case of a 10q trisomy. The only reported case of 10p trisomy was the brother of the present case (Yanagisawa and Adachi, 1970).

The case with trisomy of a whole No. 10 chromosome seemed to have more malformations (Table). A few malformations, such as harelip, cleft palate, and abnormal genitalia, were not observed in this case although they were found in cases of 10p and/or 10q trisomy. Either phenotypic variation or the presence of mosaicism could account for this. A few malformations observed in cases of 10p trisomy were not present in 10q trisomy (and vice versa). There seem to be some differences in phenotype between these two partial trisomies.

The trisomy 10 was postulated to be a clinical as well as cytotegenic entity (Nakagome et al, 1973). It may be too early to decide whether both 10p and 10q trisomy correspond to independent clinical entities. However, the phenotypic difference observed between these two conditions suggest that

![Fig. 2. Partial karyotype showing chromosomes Nos. 10 and 22 of the mother. Break points were within bands 10p11 and 22p11.](http://jmg.bmj.com/ on June 20, 2017 - Published by group.bmj.com)
this may be so. In any event, further reports of cases with these chromosome abnormalities will be of great importance in establishing these two possible new syndromes.

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REFERENCES

Stub thumbs

Summary. A case of familial brachydactyly is reported.

Bell (1951) and Temtamy and McKusick (1969) have classified the different types of brachydactyly due to maldevelopment of the phalanges or metacarpals. It may occur as an isolated phenomenon or as part of a syndrome. Type D brachydactyly or ‘stub thumbs’ is characterized by shortening and broadening of the terminal phalanges of the thumb and big toes. It is usually an isolated finding but has been reported in association with cardiac arrhythmias (Tabaznick, 1965).

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Case report

A 49-year-old manual worker was admitted to the Cardiff Royal Infirmary with a history of myocardial ischaemic pain. His stay was uneventful but he demonstrated a marked abnormality of both thumbs. They were bulbous and shortened. The nail was broad (Fig. 1). There was no abnormality of his toes. Radiology of the hands showed a shortening of the terminal phalanx in the thumbs only (Fig. 2).

His family, including reportedly the whole of the second generation, demonstrated this abnormality (Fig. 3). II.1, II.2, and II.3 had lost contact with the rest of the family and it is not known whether their progeny were affected. There were no other congenital abnormalities noted in the family.
Trisomy of the short arm of chromosome 10.

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