Three Generations and Six Family Members with a t(13q15q) Chromosome*

**Summary.** A patient with the clinical features of trisomy 13 without prosencephalic defects and with a 46,XX, 15−;t(13q15q)+ karyotype is reported. The translocation chromosome was present in five other phenotypically normal family members and could be traced back to the maternal grandfather.

The t(DqDq) Robertsonian translocation is one of the most common translocations that occurs in man (about 1:1000, Hamerton, 1971). Cohen (1971) has compiled 64 cases of t(DqDq) individuals in whom the D group chromosomes composing the translocation have been identified by autoradiography. Forty-nine of the t(DqDq) chromosomes involved numbers 13 and 14, while only six were t(13q15q), five t(13q13q), three t(14q15q), and one t(15q15q). This report presents a family with a t(13q15q) chromosome detected in six individuals and spanning three generations. The aberration was discovered when the patient described below was born and diagnosed as having trisomy D.

**Case Report**

The proposita was a premature infant, birth weight 2170 g, born to a 30-year-old gravida IV, para III, abortus I, whose prenatal course was unremarkable. Labour and spontaneous delivery were uncomplicated. There had been no history of drug intake, radiology, or illness in the mother throughout gestation. At birth (Fig. 1) the infant was noted to have an ulcerated area in midline of her scalp overlying a bony defect or third

---

Received 18 September 1972.

* This work was supported in part by a research grant (AM-02504) from the National Institute of Arthritis and Metabolic Diseases, US Public Health Service and by a grant from the Birth Defects Institute, New York State Department of Health, Albany, NY.
developed purulent discharge from the left ear from which staphylococci, resistant to penicillin, were cultured.

Her weight increased to 3277 g by the 72nd day of life, but she had difficulty with her feedings because of a poor sucking reflex. On the 73rd day of life she developed aspiration pneumonia and expired on the following day. Permission for a necropsy was denied.

Cytogenetic Studies
The proposita was thought to have an extra D group chromosome because of the clinical findings. Analysis of the chromosome complement revealed a D/D translocation chromosome, and autoradiographic studies indicated that it was composed of the long arms of chromosome numbers 13 and 15 (Fig. 2).

Chromosome analysis of the family (Fig. 3) showed the mother (II.3) to be a balanced t(13q15q) carrier. Subsequent karyotyping of other phenotypically normal family members demonstrated the balanced state in the maternal grandfather (I.1), aunt (II.2), cousin (III.1), and one sister of the proposita (III.2). The grandfather has a sister and a brother who are unavailable for chromosomal studies; both reportedly have phenotypically normal children.

Discussion
Even though the numbers of individuals in the family presented here are small, it is still interesting to compare what has occurred within this family with the findings in other t(DqDq) families. Two recent studies (Hamerton, 1968; Dutrillaux and Lejeune, 1970) have shown that there is no increase in the frequency of miscarriages among progeny of balanced male carriers. This has been the case with the present family. The maternal grandmother (I.2, see Fig. 3) experienced no miscarriages, and the grandfather (I.1) was the carrier.

There was one miscarriage in the balanced female carrier (II.3) in four pregnancies. Dutrillaux and Lejeune (1970) studied 67 families with transmissible t(DqDq) chromosomes and found that 25% of the pregnancies resulted in miscarriages when the female was the translocation carrier. An earlier study (Hamerton, 1968), however, did not find an increase in the frequency of miscarriages among the carrier females.

Hamerton (1968) also demonstrated a significant increase in balanced carriers among offspring of the male carriers; in the present case, both daughters (II.2 and II.3) of the carrier male (I.1) were balanced carriers.
Thus, the mode of inheritance of the D/D translocation chromosome found in the family reported here is quite similar to what has been demonstrated by larger studies.

RICHARD L. NEU, LYTT I. GARDNER, MARGARET L. WILLIAMS, and MATTHEW J. BARLOW*

Genetic and Endocrine Unit, Department of Pediatrics, State University of New York, Upstate Medical Center, Syracuse, New York, USA

Case Reports

REFERENCES

* Major Barlow was on assignment to the Department of Pediatrics, State University of New York, Upstate Medical Center, Syracuse, NY, from the Office of the Surgeon General, United States Army Medical Corps.

Chromosome Nomenclature

The nomenclature of human chromosomes has been standardized by a series of meetings in Denver, London, and Chicago. A report on the most recent meeting of the series which was held in Paris in September 1971 has been published* by the National Foundation of the USA. Authors are now requested to use the Paris nomenclature when submitting manuscripts. An annotation on the new nomenclature, which takes account of the banding patterns on chromosomes for the first time, will appear in the June issue of the Journal of Medical Genetics.

Three Generations and Six Family Members with a t(13q15q) Chromosome
Richard L. Neu, Lytt I. Gardner, Margaret L. Williams and Matthew J. Barlow

doi: 10.1136/jmg.10.1.94

Updated information and services can be found at:
http://jmg.bmj.com/content/10/1/94

*These include:*

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Notes**

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/