Monzygotic twins discordant for extrophy of the urinary bladder

MERETE BUGGE

From the Department of Urology A and University Institute of Clinical Genetics, Odense University Hospital, DK-5000 Odense C, Denmark

SUMMARY A pair of monzygotic male twins discordant for ectopia vesicae urinariae is described. There is no reason to think that any exogenic factors contributed to the aetiology. There was no consanguinity and no history of congenital anomalies on either side of the family. Published reports concerning family incidence and twin reports are reviewed.

Exstrophy of the urinary bladder or ectopia vesicae urinariae is a severe but rare congenital malformation. The incidence is reported to be from 1 in 30000 to 1 in 50000 births. It occurs more frequently in males than in females. The reported sex ratio varies from 2:1 to 4:1. The condition occurs in two forms: incomplete and complete. In the incomplete form a small opening corresponding to the anterior wall of the bladder is found. The complete form includes total exstrophy as well as epispadias or bifid clitoris, diastasis musculi recti abdominis, and diastasis rami superioris ossum pubis. The latter form is more frequent. Ectopia vesicae may be found with congenital malformations in almost any organic system forming complex malformation syndromes. The condition has been described in patients with Down syndrome, but only in a few cases. Extensive accounts of malformations in Down syndrome do not mention ectopia vesicae and no connection with other known chromosome anomalies has been found. The aetiology of the cases not associated with a syndrome is unknown. Nothing certain is known of the relative influence of genetic factors.

This account describes a pair of male twins who had a 99% probability of being monzygotic. One had complete ectopia vesicae and the other had no congenital malformations.

Case report

This was the first pregnancy of a 21-year-old woman. For about one year the patient had used Mini-P (norethisterone) as a contraceptive. This was first discontinued when pregnancy was ascertained 5 weeks after the last normal menstruation. In the first 5 weeks of pregnancy the patient had taken Duspatalin tablets (Mebeverini chloridum), at the most three tablets in all. In the first two months of pregnancy she had an almost constant brown discharge per vaginam and twice had minor spotting. At approximately the eighth week of pregnancy the patient was admitted to a department of gynaecology for imminent abortion after heavy unprovoked fresh vaginal bleeding. The condition subsided after a period of rest in bed, and the pregnancy continued without complications. The mother was in good health during the entire pregnancy and in particular did not suffer from any febrile disease. She was at no time exposed to known toxic agents at her place of work. Abdominal x-ray at 33 weeks' gestation confirmed a suspected twin pregnancy. X-ray examination was not performed earlier in the pregnancy, and at no point was ultrasound examination carried out. Spontaneous and uncomplicated delivery occurred in the 38th week of gestation in an obstetric department. Both twins were delivered by the vertex.

TWIN A
This was a live boy, birthweight 2850 g. Immediately after birth it was observed that the infant had complete ectopia vesicae. Intravenous urography showed an otherwise normal urinary system and there were no other notable malformations. Surgery was carried out, and by the age of 3 years the child was normally developed for his age, height 94·5 cm, weight 13·5 kg.

TWIN B
This was a live boy, birthweight 3250 g. No sign of
congenital malformation was found in this infant. The child was normally developed at the age of 3 years and had never suffered from any serious illness. His height was 97.5 cm, and weight was 14.5 kg. X-ray examination of the urinary tract was not considered to be justified as the boy had no symptoms.

The parents were in the same age group and both were healthy and unrelated. On examination of the family history no evidence was found of this or other congenital malformations in any of the close relatives.

At birth the placenta was examined by an experienced obstetrician, who found a fused placenta with monochorionic, biamniotic membranes. At the age of 3 years the children looked almost completely alike. They have the same colour eyes and hair. Blood group determinations, carried out when the children were aged 2 years, showed total identity in 18 serological systems. The systems examined included the ABO, MN, Rh (5 antigens), P, Kell and Duffy blood groups, the Hp, Gc, Gm (a,x,b) and Inv serum protein groups, and the PGMI, AcP, GPT, EsD, GLO, AK, PGD, and ADA enzyme polymorphisms (The Institute of Forensic Medicine, The University of Copenhagen).

These findings, with a probability of more than 99%, justify the diagnosis of monozygosity.

Discussion

In the first weeks of pregnancy the mother ingested Duspatalin (Mebeverini chloridum). There are no published reports of congenital malformations occurring in children of mothers who ingested this medication during pregnancy. The mother continued her peroral contraceptive up to 5 weeks after her last normal menstruation. Congenital malformations associated with the urogenital system resulting from the ingestion of norethisterone during pregnancy have been described, but even the more extensive accounts do not mention ectopia vesicae.

Familial cases of ectopia vesicae are described in published reports, but the descriptions are few and incomplete. In his work in 1958 Higgins reports having treated 147 children with the condition. He mentions briefly having seen the malformation occur twice in "members of the same family". Further details, however, are not given. In 1928 Kohler described a boy whose male cousin had the same malformation. Four publications report bladder extrophy in two or more sibs. Glaser and Lewis described a healthy woman whose first and second child, both boys, born within a 2-year period, had ectopia vesicae. He mentions that there was no known familial predisposition to this or other congenital malformations. Chrisholm briefly mentions having treated a boy whose sister had the same complaint. Schilling describes a similar case just as briefly. Randall and Harwick treated a girl surgically for ectopia vesicae and mentioned that her two brothers had the same malformation. She herself gave birth to a healthy child, and there are no accounts of patients with ectopia vesicae giving birth to children with the same condition. A number of authors emphasise that they have never experienced such instances.

Four pairs of twins with ectopia vesicae are described in published reports. One discordant pair was of different sex and therefore certainly dizygotic. The twins in the three other pairs were of the same sex and included two pairs of boys, who were concordant, and one pair of girls who were discordant. One of the male pairs were described in 1943 by Higgins as "identical", but no mention was made of the reasons for this diagnosis. Only one placenta was found. He also mentioned that no similar cases were found in the family. Smith and Lattimar described the other pair of boys. These were also called "identical", but no further information was given. Coates described the only pair of girls. The description did not indicate any mention of observations made for the purpose of establishing the zygosity of the twins. Thus, none of these reports contains definite information about the diagnosis of zygosity and no definite conclusion can be reached regarding possible genetic aetiology from the information given.

The present case in which ectopia vesicae occurred discordantly in a monozygotic twin pair allows the conclusion that the condition is at least not purely genetically determined. The present report does not allow for any conclusions regarding the aetiology. The limited information available suggests that an increased recurrence risk, albeit low, among sibs of affected subjects may exist. It seems, however, desirable to have some estimate of this risk before recommending fetoscopy to women who have had one child with extrophy of the urinary bladder. As far as the author is aware there is no mention in published reports of the malformation not associated with a syndrome diagnosed prenatally by ultrasound examination or amniocentesis.

References

Monozygotic twins discordant for exstrophy of the urinary bladder


Requests for reprints to Dr M Bugge, Department of Urology A, Odense University Hospital, DK-5000 Odense C, Denmark.