was noted (fig 1). Bilateral postaxial polydactyly was found in both upper and lower limbs (fig 2). External genitalia were at an undifferentiated stage. There were only two vessels in the cord. The whole body was serially sectioned, stained with haematoxylin and eosin, and examined microscopically. Internal anomalies included alobar holoprosencephaly with absence of midline structures, ventricular septal defect (membranous portion), and single umbilical artery. No abnormalities were noted in the trachea, lungs, gastrointestinal tract, liver, pancreas, or urogenital organs. Histologically, the gonadal sex was male.

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Syndactyly, ectodermal dysplasia, and cleft lip and palate

SIR,

Recently, each of us reported independently in the journal a family in which two children were affected with a syndrome which seemed to be new.^{1,2}

TABLE Main clinical features of the syndrome.

	Zlotogora et al ¹		Ogur and Yukse ²	
Sex	Male	Female	Male	Male
Cleft lip and palate	+	+	+	+
Partial syndactyly				
Fingers	3-4	3-4	2-3-4	2-3-4
Toes	2-3*	2-3*	2-3	2-3
Ectodermal dysplasia				
Abnormal hair	+	+	+	+
Pili torti	+	-	+	+
Thickened, dry skin	+	NR	+	+
Abnormal teeth	+	NR	+	+
Normal nails	+	+	+	+
Mental retardation	Mild‡	NR		Moderate
Consanguineous parents		+		+

- = Not present in the child. NR = not relevant since the child died very young.

*Owing to an error in the original article the syndactyly of the toes was reported to be between toes 3 and 4.

‡Appeared at the age of four years, mainly on the palms and soles.

§At the age of four years the child had caught up most of the delay and was only mildly retarded. Speech was very delayed.

Comparison of the clinical signs show that although the families are unrelated, the affected children present with very similar features, which are summarised in the table. It seems that this represents a distinct syndrome which is inherited as an autosomal recessive disorder. Additional cases are needed for further delineation of the syndrome.

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Are 'upper' and 'lower' neural tube defects aetiologically different?

SIR,

The idea that anencephaly and spina bifida cystica are aetiologically related, since each occurs with increased frequency in sibs of probands with the other,¹ may need reconsideration. In two studies,^{2,3} families of probands with neural tube defects were classified according to whether the lesion was 'upper' (anencephaly and thoracic spina bifida, which arise by failure of neurulation) or 'lower' (lumbar and sacral, which represent errors in canalisation⁴). All of 25 sib pairs were concordant as to level, suggesting that the two types are genetically different.^{2,3} T11 to T12 was considered the dividing point between upper and lower lesions.

In contrast, during an epidemiological study of neural tube defects in Newfoundland,⁵ we noted 11 pairs of affected sibs, without other malformations, of which four were discordant for level of lesion.

In family A, a girl was born with a lumbosacral myelomeningocele involving L1 to S3, with diastematomyelia of L1 (lower), followed by a sister with anencephaly and cervical spina bifida (upper). In family B, a girl with a lumbosacral myelomeningocele involving L1 to S3 (lower) was followed by a brother with a myelomeningocele of T10 to S3 (upper). In family C, the first boy was born with a tuft of hair in