In 1971 Johanson and Blizzard\(^1\) reported a new syndrome in three unrelated girls characterised by congenital aplasia of the alae nasi, deafness, hypothyroidism, dwarfism, absent permanent teeth, and malabsorption. Children with this syndrome had been described earlier by Morris and Fisher in 1967\(^2\) and Townes in 1969\(^3\) as examples of trypsinogen deficiency disease. Townes and White\(^4\) subsequently reviewed the patient reported in 1969\(^3\) and described the presence of additional features which confirmed the diagnosis of the Johanson-Blizzard syndrome. There have since been 22 patients reported with Johanson-Blizzard syndrome, and a further seven children related to these. The spectrum of associated features is now well documented and the inheritance of the syndrome is autosomal recessive. However, there remain many problems which make counselling difficult, in particular the degree of mental handicap and the observation that some children die from complications of the severe malabsorption despite intensive treatment. This article reviews the 22 patients previously reported and also includes details of a previously unreported boy.

**Clinical features**

The main features are shown in the table. The most constant signs necessary to make a diagnosis are aplasia of the alae nasi, an exocrine pancreatic defect, and unusual hair growth pattern.

In the absence of major structural abnormalities, the affected infant usually comes to medical attention because of failure to thrive. On presentation, the infant is malnourished, hypotonic, and often oedematous because of hypoproteinaemia.

**THE FACE**

The diagnosis should be made easily in the neonatal period. Indeed the 'gestalt' of the Johanson-Blizzard syndrome is so distinct that seen once it should not be missed (figs 1 and 2). The picture is determined by the unusual nasal configuration. The severe aplasia of the alae nasi leads to the appearance of a

### Table: Frequency of features in the Johanson-Blizzard syndrome

<table>
<thead>
<tr>
<th>Feature</th>
<th>Reported cases (n=22)</th>
<th>This case</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypoplastic alae nasi</td>
<td>22</td>
<td>+</td>
</tr>
<tr>
<td>Pancreatic insufficiency/failure to thrive</td>
<td>21</td>
<td>+</td>
</tr>
<tr>
<td>Aplasia cutis congenita</td>
<td>19</td>
<td>-</td>
</tr>
<tr>
<td>Short stature</td>
<td>15</td>
<td>+</td>
</tr>
<tr>
<td>Mental retardation</td>
<td>13</td>
<td>Mild</td>
</tr>
<tr>
<td>Dental anomalies</td>
<td>13</td>
<td>+</td>
</tr>
<tr>
<td>Deafness</td>
<td>12</td>
<td>+</td>
</tr>
<tr>
<td>Anorectal anomalies</td>
<td>11</td>
<td>-</td>
</tr>
<tr>
<td>Microcephaly</td>
<td>8</td>
<td>-</td>
</tr>
<tr>
<td>Genitourinary abnormalities</td>
<td>7</td>
<td>-</td>
</tr>
<tr>
<td>Hypothyroidism</td>
<td>6</td>
<td>+</td>
</tr>
<tr>
<td>Cardiac malformation</td>
<td>3</td>
<td>-</td>
</tr>
</tbody>
</table>

**FIG 1** Our patient, a boy, aged two months. Note the small nose with aplasia of the alae nasi and the abnormal hair growth pattern, also shown in fig 3.
thin, torpedo shaped nose with large nostrils. In addition, the hair is swept up, especially frontally (fig 3), and has a patchy distribution over the scalp. Closer inspection shows areas of alopecia (fig 4) with underlying aplasia cutis congenita, which are characteristically in the midline and in the occipital region. They heal to form atrophic scars.

MALABSORPTION
An exocrine pancreatic defect is a constant feature of this condition. Townes3 and Townes and White4 described the abnormalities of pancreatic function. They reported an absence of trypsin, chymotrypsin, and their proenzymes, as well as carboxypeptidase and lipase, but they thought the amylase activity was normal. This was subsequently found to be a spurious result as isoenzyme studies showed the amylase was from the salivary gland. When the contribution from saliva is excluded, pancreatic amylase activity is absent. At necropsy the parenchyma of the pancreatic gland is replaced by fatty tissue.5 6 There are fewer islets of Langerhans but clinically there are no reports of impaired glucose tolerance.

The severe malabsorption caused by these enzyme deficiencies leads to hypoproteinaemia, oedema, anaemia, and failure to thrive. The treatment of the pancreatic failure is by pancreatic enzyme replacement, as in cystic fibrosis. It is a life threatening condition and several of the children reported have died despite full medical treatment.

SHORT STATURE
The short stature has been attributed to hypothyroidism and malabsorption. There have been su...
Syndrome of the month

Syndrome cases cause. The stature short has yet thyroid to may difficult at test had normal here were mention no as genital, as hypothyroidism may at thyroid function. Reduced HYPOTHYROIDISM At mU/l) have tion dren however, is developmentally is brain to confirmed as mental retardation. However, this cannot have also been considered in terms of the hypothryoidism. However, there are sufficient children with normal thyroid function to know that hypothyroidism is unlikely to be the cause, though it can contribute if allowed to go untreated.

MENTAL RETARDATION
Thirteen out of the 22 children reported have been developmentally delayed. This is an underestimate as infants have died in the neonatal period. The cause is obscure. Daentl et al.6 showed focal migrational defects in the brain at necropsy but this was not confirmed by Moechsler et al.,8 who found the brain to be small but structurally normal. There is also no close relationship between mental retardation and hypothyroidism. As pointed out by Moechsler and Lubinsky,6 the patients of Schussheim et al.5 and Sismansis et al.7 were severely retarded but neither microcephalic nor hypothyroid. The degree of retardation cannot be predicted. The patient of Day and Israel10 showed mild developmental delay and that of Townes11 was reported as “relatively normal” at three and a half years, although by 12½ years she needed special education. The brother and sister reported by Moechsler and Lubinsky8 were normal at two and a half and two years. Severe retardation was found in the patients of Daentl et al.,5 Baraitser and Hodgson,12 and Mardini et al.13

ANORECTAL ANOMALIES
As shown in the table, 11 of the reported children with the Johanson-Blizzard syndrome had anorectal abnormalities. In the majority of cases this was an imperforate anus. These children come to medical attention early and the initial surgical management is the fashioning of a transverse colostomy. It is important that poor weight gain in these infants is not attributed to the surgery but that malabsorption is recognised and treated.

DEAFNESS
Hearing loss has been reported in 12 out of 22 patients. The case of Sismansis et al.5 was investigated in detail. There was a severe sensorineural hearing loss with associated absent vestibular function, but the inner ears were structurally normal on polytomograms. There have been no further reports of more detailed radiological investigations. At necropsy6 the temporal bones were not studied.
Our patient has a symmetrical, moderately severe sensorineural hearing loss.

OTHER FEATURES
Additional features listed in the table are abnormalities of dentition, genitourinary anomalies, and cardiac malformations. The dental findings have been well reviewed by Zerres and Holtgrave.14 The children have delayed eruption of teeth, which are small but normal in shape. The genitourinary anomalies were striking in the three girls originally reported,1 two of whom had a single urogenital orifice, but other children have not had major structural problems. A congenital heart defect has been reported in only three of the children, two of whom were sibs reported by Helin and Jodal15 with situs inversus. Minor abnormalities of the lacrimal duct have also been recorded.

DIFFERENTIAL DIAGNOSIS
The diagnosis of the Johanson-Blizzard syndrome is not difficult when all the features are present. Hypoplasia of the alae nasi occurs in the ocudontodigital syndrome,16 aplasia cutis congenita of the scalp in the Adams-Oliver syndrome,17 and pancreatic malabsorption in the Shwachman-Diamond syndrome, but these should not prove to be diagnostic problems.
Counselling

There is strong evidence for autosomal recessive inheritance. Affected sibs have been described by Moeschler and Lubinsky,8 Day and Israel,10 Helin and Jodal,15 and Bresson et al.18 Parental consanguinity has been reported by Schussheim et al.,9 Sismanis et al.,7 Mardini et al.,13 and Bresson et al.18

A more difficult problem in counselling is that of predicting the degree of mental retardation, the presence of severe structural lesions, and the success in treating the pancreatic exocrine defect. There have been no reports of prenatal diagnosis to date. At present one could offer a high resolution ultrasound scan in the hope of detecting the distinctive facial features and any structural abnormalities, such as the cardiac or urogenital lesions.

Prognosis

In the family of Mardini et al.,13 all three patients died in infancy from complications of malabsorption and failure to thrive. The girl reported by Townes3 and Townes and White4 was still alive at the age of 12 years nine months. She had short stature, no permanent teeth, a sigmoidostomy because of an imperforate anus, and she required pancreatic enzyme supplements with her meals. She had mild mental retardation. If the pancreatic malabsorption problems are overcome the child can survive infancy, but is likely to require prolonged medical supervision. Even when given the best medical attention these children may develop severe problems associated with hypoproteinaemia, namely infections and oedema, which can lead to death in childhood.

References

17 Adams FH, Oliver CP. Hereditary deformities in man due to arrested development. J Hered 1945;36:3-7.

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