Conductive Hearing Loss and Malformed Low-set Ears, as a Possible Recessive Syndrome*

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More than 50 phenotypically distinct forms of hereditary deafness have been found in man (Konigsmark and McKusick, 1966; Procter and Procter, 1967). On the basis of associated defects caused by the same gene, these can be separated into 10 different groups. These groups include: deafness associated with integumentary system disease (Robinson, Miller, and Bensimon, 1962), heart disease (Fraser, Froggatt, and James, 1964), skeletal disease (Forney, Robinson, and Pascoe, 1966), eye disease (Alström et al., 1959), nervous system disease (Matthews, 1950), renal disease (Braun and Bayer, 1962), thyroid disease (Batsakis and Nishiyama, 1962), external ear malformations (Fourman and Fourman, 1955; Wildervanck, 1962), as well as deafness with no associated defects, and hereditary deafness with other associations.

A Mennonite group has been found to offer an excellent opportunity for the study of some recessive hereditary diseases, for two reasons: excellent family records are available and the rate of consanguineous marriages is high.

The two interrelated families described here were found in a health survey of conservative Mennonites in Lancaster County, Pennsylvania. In Family I, 2 of 8 children were affected with malformed low-set ears and conductive hearing loss, while in Family II, 3 of 12 living sibs were similarly affected. Though two types of external ear deformities and hearing loss have been reported (Fourman and Fourman, 1955; Wildervanck, 1962), the deformities are quite different from those in the two branches of this family. The affected children recorded here probably have a heretofore undescribed type of familial hearing loss, recessive in transmission, and characterized by malformed low-set ears with conductive hearing loss.

Material and Methods

A pedigree was obtained by information from each family and by records in the Mennonite Information Center.

Each of the affected children was tested in the Hospital's Hearing and Speech Center, with a standard testing procedure and criteria (Berlin, 1965). The battery of tests included: air conduction, bone conduction, speech reception threshold, speech discrimination and tone decay; the short-increment sensitivity index (SISI), and the alternate binaural loudness balance test were done when appropriate. In addition, all affected children had caloric vestibular tests, and physical and neurological examinations. Chromosomal analyses were done on two affected children, skull X-rays on two, temporal bone polytomograms on one, and an exploratory tympanotomy on one patient. Psychological testing was done on three affected children.

Findings

Ear Deformity. All six affected children showed this defect to some extent bilaterally. The pinna was small, sometimes low-set, with a large overturned skin flap at its dorsal edge. The mildest form of the deformity showed only minimally deformed pinna. However, in the most severe forms, the pinna was low-set, quite small, and half-moon in shape (Fig. 1, 2, and 3).

Mental Status. By history three of the six affected children (Fig. 4) had been slow in sitting, walking, and speaking. One (II. 1) did not walk until 5 years of age. Now, at the age of 19, he can feed and dress himself and do simple household tasks.

Intelligence tests on four of the affected (I. 1, II. 1, II. 3, II. 4; Fig. 4) placed three (I. 1, II. 1, II.
Hearing. Pure tone audiograms showed four of the six affected children to have a 70 to 80 dB HL ISO three frequency average loss in at least one ear. Bone conduction testing showed minimally decreased hearing, indicating a conductive loss with pathology in the middle ear. SISI and recruitment tests were negative, indicating no cochlear damage. Tone decay tests were also negative, suggesting no retrocochlear pathology. Speech reception threshold scores validated the pure tone audiometry. In one patient (II. 1), we were unable to get any response to tones to 110 dB HL ISO. This patient was severely retarded and could not understand test instructions. Another child (II. 4) could not be tested because of low IQ, giving erratic responses at 80 to 100 dB HL.

Ossicular Chain Deformity. An exploratory tympanotomy, was performed on one (II. 2). At operation, a single ossicle was noted; this had the shape of a malleus and was somewhat posteriorly positioned. No incus was seen. A loose cord, attached to the head of the malleus, passed medially to the oval window area. Inspection of the oval window revealed no foot-plate or superstructure of the stapes present. The promontory was relatively normal, and a round window niche could not be seen. The bone in the area of the oval window was trephined, removing approximately 2 mm. of bone, leaving in a very thin plate of bone over the vestibule. This plate was then removed, and a slight leak of perilymph noted. A No. 6 Teflon prosthesis was placed into the opening, and its wire end was attached to the neck of the malleus. The prosthesis was noted to move with the malleus when this was manipulated.

The patient was noted to have a 10 to 15 dB improvement in hearing on subsequent auditory tests, but because his other ear had a less severe hearing loss, this improvement was not clinically apparent.

Physical and Neurological Examination

Small Stature. Three of the six affected children were somewhat smaller than the average of their sibs for that age.

Head and Neck. In addition to the ear deformity, three of the six affected children had a high arched palate, not noted in any of their non-affected sibs. Otherwise, the head and neck examination showed no abnormalities.

Cardiovascular. Five of the six affected children had a grade 1-4/6 systolic blowing murmur heard best at the cardiac apex. None of the unaffected sibs had such a murmur. Electrocardiograms were normal showing no cardiomegaly or axis deviation. There were no symptoms of cardiovascular disease.
Other Findings. Two of the four affected males (I. 2 and II. 1) had cryptorchidism with no testes palpable in the scrotum or canals. These and one other (II. 2) had hypogonadism, but chromosome examinations done on two of these (I. 2 and II. 2) were normal.

The two children in Family I had genu valgum, but this may be an independent trait since their mother had a similar appearance and none of the children in Family II were affected. These two children also showed slight clinodactyly, not seen in any of the other children.

One child (II. 1) had marked pes planus and valgus, not seen in any of the other children.

Neurological and vestibular examinations were normal in all six affected children.

Case Reports

Case 1. The proband (I. 1) was a 10-year-old girl, with one ear mildly affected and one moderately affected. She first sat up at 9 months, later than her 2 older sibs, though she did walk at 13 months. The family gradually became aware of a hearing loss, probably alerted to expect it by the deformed external ear. She spoke her first words at about 2 years of age, and could form sentences by 6 years of age. She did well in a special class in public school. Testing at her school placed her IQ in the 40 to 60 range.

On physical examination she was small for her age (117 cm., 22.3 kg.), with a grade 2/6 systolic ejection murmur loudest at the apex, and transmitted to the carotids. She had slight asymmetry of the mouth and bilateral genu valgus. Physical examination was otherwise normal. Audiologically, her right ear was normal;
Conductive Hearing Loss and Malformed Low-set Ears, as a Possible Recessive Syndrome

Case 2. I. 2, her brother, was 6 years old and had more severely affected pinnae bilaterally than his older sister. His father felt he had been slow in development, recalling only that he walked at about 14 to 16 months. He attended the local parochial school, doing fair work. He was a pleasant boy, slightly small for his age (113.6 cm., 21.3 kg.). He did not speak sentences but understood and obeyed very well. His mental retardation was milder than that of his sister.

Physical examination showed a harsh blowing systolic murmur (Grade 4/6) loudest at the apex. His reflexes were brisk but symmetrical. He had extremely small genitalia, with well-marked median raphe and no testis in his scrotum. Chromosome cultures showed a normal karyotype. He had slight genu valgum. Audiological testing showed a severe conductive deafness bilaterally in the left ear showed a conductive hearing loss of 60 to 70 dB HL ISO (Fig. 5), with 50 to 60 dB narrow band masking in the appropriate ear.
**Case 4.** II. 2, an 18-year-old brother, was 166.4 cm. tall and weighed 56.9 kg. He spoke poorly, probably due to poor hearing. He was able to finish the eighth grade at a parochial school in regular classes. His pinnae were severely affected (Fig. 2 and 3); one was quite low-set.

**Case 5.** II. 3, a 6-year-old boy, was normal on physical, neurological, and vestibular examination except for a 2/6 systolic murmur. The best two frequency averages in the speech range were 67 dB HL in the left ear and 5 dB HL in the right ear. His pinnae were mildly deformed (Fig. 2 and 3).

He had a mental age of 5 8/12 years with an IQ of 98 on the Stanford Binet intelligence scale Form L-M. He scored in the mental age level 3–9, with a quotient of 65 on the Peabody Picture Vocabulary Test. His vocabulary was poor relative to his normal intelligence.

**Case 6.** II. 4, a 5-year-old girl, had a completely normal physical, neurological, and vestibular test. She had no systolic murmur. Her language development score was below the 18-month level, with visual motor and form perception functioning in the 2- to 3-year range. Her over-all ability was in the 40 to 60 IQ range. Though she apparently could hear well in response to verbal commands, we were unable to obtain criterion level pure tone test results with this child.

**Remainder of Family.** The parents in these two families were not affected with the syndrome described. In Family I, the maternal great grandfather and two paternal uncles had a hearing loss in their 5th to 6th decade of life. In Family II, the father was 46 years old and had a neural hearing loss without an air-bone gap. The paternal grandfather had a hearing loss since his 6th decade of life, and three paternal cousins were mentally retarded, one with histidinaemia.

**Discussion**

**Syndrome.** The features of this syndrome can be divided into those which appear in all affected and those which are inconstant. Features present in all affected include: (1) pinna deformity, varying greatly in degree (Fig. 1, 2, and 3), (2) conductive hearing loss, usually bilateral, and varying from mild to severe. The following features were present in only some of the affected, but probably are features of the syndrome: (a) low-set ears, present in three of the six affected; (b) mental retardation, present in three of the affected; (c) immature external genitalia, present in all affected males.

In addition to these features, five of the six affected children were smaller than their sibs. Three had a high arched palate, and five had a systolic murmur with normal electrocardiograms and
Conductive Hearing Loss and Malformed Low-set Ears, as a Possible Recessive Syndrome

Fig. 5. Audiograms of all affected children.

Air conduction
Bone conduction (with appropriate ear masked)

Right
Left
Mengel, Konigsmark, Berlin, and McKusick

TABLE
SUMMARY OF MAJOR CLINICAL FINDINGS IN EACH OF AFFECTED CHILDREN

<table>
<thead>
<tr>
<th>Age (yr.)</th>
<th>I. 1</th>
<th>I. 2</th>
<th>II. 1</th>
<th>II. 2</th>
<th>II. 3</th>
<th>II. 4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td>F</td>
<td>M</td>
<td>M</td>
<td>M</td>
<td>M</td>
<td>M</td>
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<td>Right ear deformity</td>
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<td>+</td>
<td>+++++</td>
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<td>+++++</td>
<td>+++++</td>
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<tr>
<td>Right ear hearing loss*</td>
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<td>55</td>
<td>No response</td>
<td>+55</td>
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<td>+55</td>
</tr>
<tr>
<td>Left ear hearing loss*</td>
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<td>82</td>
<td>No response</td>
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<td>77</td>
<td>77</td>
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<tr>
<td>Mental retardation</td>
<td>+++++</td>
<td>0</td>
<td>+++++</td>
<td>0</td>
<td>0</td>
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Key: 0—not present, +—mild, + +—moderate, + + +—severe defect.
* 3 frequency average in dB HL ISO.

Origin of Hearing Loss. Audiograms on most of the affected children indicated a conductive hearing loss. In one patient (II. 2) the hearing loss was due to malformation of the middle ear bones, as documented at operation. Probably, middle ear malformation is responsible for the hearing loss in the remaining affected children.

Type of Transmission. The family trees were traced for 12 generations. All four parents were descendants of a man who died in Switzerland in 1720. His four sons came to this country in the early 18th century. It appears that this syndrome is transmitted as a single autosomal recessive gene and was carried to this country by at least two sons of the progenitor.

Treatment. The mild improvement in hearing in one of our patients (II. 2) by insertion of a prosthesis suggests that patients with this affection deserve a complete otological and audiological study, with the aim of possible surgical correction of the hearing loss.

Summary

In a health survey of a genetic isolate, six cases of pinna deformity and conductive deafness were found. Tympanoplasty in one patient showed malformation of the middle ear bones. Other features, including mental retardation and hypogonadism, may be part of the disease. The mode of transmission is autosomal recessive.
We wish to thank Dr. Minerva Stauffer for bringing these families to our attention.


REFERENCES


