A syndrome of insulin resistance resembling leprechaunism in five sibs of consanguineous parents

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Abstract

Leprechaunism is a rare autosomal recessive disorder associated with extreme insulin resistance with paradoxical hypoglycaemia. It is characterised by prenatal and postnatal growth retardation, reduced subcutaneous tissue, coarse features, acanthosis nigricans, enlarged genitalia, and death in the first year of life. Defects in both the insulin receptor and postreceptor steps of the insulin action pathway have been reported. At the molecular level, several mutations have been described.

The patients reported here are from a Yemeni family with a syndrome of insulin resistance similar to leprechaunism in which the parents are second cousins and five of their eight children are affected. However, the phenotypes seem to be less severe than the classical leprechaunism previously described. All the children are alive (oldest 11 years), there is normal subcutaneous tissue, and a normal growth pattern in some of them. It may be that this is a milder type of leprechaunism with a better prognosis, perhaps caused by a different type of mutation from those previously described.

(J Med Genet 1993;30:470-5)

Leprechaunism is a rare autosomal recessive disorder characterised clinically by intrauterine growth retardation, diminished fat and muscle tissue, characteristic facies, acanthosis nigricans, enlarged genitalia, and early death. Biochemical abnormalities include hyperinsulinaemia, insulin resistance, and impaired glucose homeostasis.

This syndrome was originally described by Donohue in 1948 who reported a female infant looking like an elf with multiple endocrine abnormalities. This was presented as a case of 'dysembryocrinism'. In 1994 a similar child was born to the same family and Donohue and Uchida proposed the term 'leprechaunism' for this condition.

In the consanguineous Yemeni family reported here five children are affected by a syndrome of insulin resistance resembling leprechaunism, but the disorder seems to be less severe than the classical previously described type.

Family report

The family consists of second cousin parents (mother 28 years and father 33 years) of Yemini origin with eight children, five of whom have the disorder, four males and one female (fig 1). Both parents appear normal.

CASE 1

The proband (IV-3), a boy, was born in 1983 after a normal pregnancy and delivery. His birth weight was 2500 g. He had convulsions immediately after birth and was found to have hypoglycaemia with a fasting blood sugar of 0.8 mmol.l⁻¹ and a serum insulin level of 247 μU.ml⁻¹. He was treated with frequent feeding, prednisolone, and later diazoxide but with no response. At the age of 7 months he underwent subtotal pancreatectomy. Histopathology disclosed islet cell hyperplasia with no signs of nesidioblastosis. His therapy remained as before. Nevertheless at 9 months of age further pancreatic excision was done and only a small crescent of pancreatic tissue was left. He continued, however, to have symptomatic hypoglycaemia and his insulin level remained high at 1000 μU.ml⁻¹. At 4 years of age he spent three days on a ventilator because of coma and seizures. At the age of 5 years he developed a full blown picture of diabetes mellitus with polyuria and polydipsia. His blood sugar was 27 mmol.l⁻¹ with a simultaneous insulin level of 1000 μU.ml⁻¹. He is 7 years old now and on no medication. He still has frequent symptomatic hypoglycaemia and is moderately mentally retarded. His weight is 17 kg, height 112 cm, and head circumference 48 cm, all below the 5th centile. He has the following dysmorphic features (fig 2A, table 1): widely spaced eyes, depressed nasal bridge with flared alae nasi, thick lips, large ears, gum hypertrophy, prominent nipples, and distended abdomen with large genitalia (fig 2B). He has acanthosis nigricans at the back of the neck, in both axillae, and in the abdomen and groin (fig 2C). There is generalised hypertrichosis, scattered white scalp hair (fig 2D). The toe nails are short and convex. He has large hands and feet with brachydactyly.
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There is no reduction in subcutaneous fat and muscle.

Auscultation indicated a loud systolic murmur (grade V) at the left sternal edge, and echocardiography showed thickened myocardial muscles of the left and right ventricles. Renal ultrasound showed bilaterally enlarged kidneys (right 9.1 cm, left 10.4 cm). CT scan of the brain and EEG were normal. Bone age corresponded to chronological age. Chromosome analysis showed an apparently normal female karyotype (46,XX). Her lowest blood sugar recorded was 0.7 mmol.l⁻¹ with a fasting insulin level of 131 μU.ml⁻¹. The highest recorded insulin level was 1050 μU.ml⁻¹.

CASE 3
A boy (IV-1) was born in 1980 by normal delivery after a normal term pregnancy. The birth weight was 2800 g. Soon after birth he started to have hypoglycaemic attacks during which he went limp and blue and both eyes rolled up. No specific treatment was given and the mother was able to control these attacks with frequent feeding. He is 11 years old now and still gets attacks of abnormal behaviour if not fed frequently. He is, however, attending school and is doing well at the appropriate age grading.

Examination at the age of 11 years showed a weight of 30 kg, height of 137 cm (both on the 25th centile), and head circumference of 53 cm. He has a similar phenotype to his other sibs (fig 4A–C, table 1).

Echodardiography showed moderate thickening of the left and right ventricular muscles. Renal ultrasound showed slightly enlarged kidneys (right 10.2 cm, left 10.2 cm).

Relevant laboratory investigations were: fasting blood sugar of 2.2 mmol.l⁻¹. The lowest and highest levels of insulin were 74 μU.ml⁻¹ and 1690 μU.ml⁻¹, respectively.

CASE 4
A boy (IV-5) was born in 1986 after an uneventful pregnancy by normal vaginal delivery. His birth weight was 2500 g. He appeared normal until the age of 2 years when he started to have attacks of sweating and loss of vision after prolonged fasting. The parents have managed these attacks by frequent feeding. He has normal developmental milestones.

Examination at the age of 5 years showed a weight of 19 kg and a height of 108 cm, on the 25th and 50th centiles respectively. His head circumference is 49 cm. He has a very similar phenotype to his other sibs (fig 5, table 1).

Echodardiography showed thickened myo-
cardiac muscles of the left and right ventricles. Renal ultrasound showed bilaterally enlarged kidneys (right 4.9 cm, left 10 cm). Laboratory investigations showed a blood sugar of 1.3 mmol.l⁻¹ and insulin level of 656 μU.ml⁻¹.

**CASE 5**
A boy (IV-8) was born in 1990 by normal vaginal delivery after an uneventful term pregnancy. His birth weight was 2600 g. He was found to have the same phenotype as his other sibs (fig 6, table 1), and because the mother was aware of the problem she started feeding him frequently. As a result he did not develop symptomatic hypoglycaemia. He was admitted to hospital on a few occasions because of asthmatic bronchiolitis and gastroenteritis.

He is now 1 year 7 months old with a weight of 8 kg, height of 73 cm, and head circumference of 48 cm. He has normal developmental milestones.

Echocardiography showed markedly hypertrophied right and left ventricular muscle suggestive of hypertrophic cardiomyopathy. Renal ultrasound showed an enlarged left kidney (8.1 cm) and a normal right kidney. Bone age corresponded to chronological age.

Relevant laboratory results included blood sugar of 2 mmol.l⁻¹, insulin level 64 μU.ml⁻¹ (fasting), and 845 μU.ml⁻¹ (postprandial).

In all five sibs serum levels of glucagon, cortisol, somatotropin C, FSH, LH, T3, T4, TSH, and 17-hydroxyprogesterone were within normal limits.

**Discussion**

Elsas et al summarised the phenotype of leprechaunism as follows: severe intrauterine growth retardation, small elfin face with protruberant ears, distended abdomen, relatively large hands, feet, and genitalia, and abnormal skin with hypertrichosis, acanthosis nigricans, and decreased subcutaneous fat. The majority of the 52 previously reported patients share most of these typical features of leprechaunism but there are others who do not and in these the diagnosis has been questioned.

Patterson and Watkin described a probable case in a male patient, but later Patterson suggested that this patient might have had a different disorder, since there were clinical signs of 'Cushing's disease' with enlarged adrenals and severe changes in the bones at necropsy. David and Goodman reported similar observations under the name 'Patterson syndrome'. Dallaire et al described three infants with generalised elastic fibre deficiency and leprechaunoid features. The majority (61.1%) of the cases in whom the diagnosis was unequivocal died in the first year of life, 90% of cases had loss of subcutaneous fat, 41% had poor weight gain after birth, 42% of cases had intrauterine growth retardation, and 38% of cases had mental retardation (table 1).

The five children in this report have many of the features of leprechaunism (table 1) but seem to be less severely affected than the previously reported cases. They are all alive and are mentally normal apart from one patient (IV-3) whose mental deficiency could be attributed to prolonged hypoglycaemic attacks. None has any reduction in subcutaneous fat. The hypoglycaemia seen in all these children is easily controlled by frequent feeding. Three have mild growth retardation (IV-3, IV-7, IV-8) and two have normal growth patterns (IV-1, IV-5), but none suffered intrauterine growth retardation.

However, all five children have kidney
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enlargement on ultrasound (table 1). This finding has been reported in only five patients with leprechaunism but, since most were found at necropsy, it may be that kidney enlargement had not been suspected in the other patients reported. Histological examination in those patients showed tubular dilatation or calcification with intratubular granules or both. Ellis et al, however, reported a 13 year old leprechaun child with hyperglycaemia and hyperinsulinaemia who had hypertension, microalbuminuria, and enlarged kidneys. The findings on renal biopsy were similar to those seen in diabetic nephropathy.

Other genetic syndromes with insulin resistance include type A insulin resistance, lipoatrophic diabetes mellitus, and Rabson-Mendenhall's syndrome. Although these syndromes share some features in common (for example, acanthosis nigricans and hyperandrogenism), a syndrome can be defined based on the presence or absence of specific clinical features. For example, in lipoatrophic diabetes there is atrophy of subcutaneous fat and hypertriglyceridaemia. Rabson-Mendall's syndrome is associated with abnormalities of the teeth and nails and reportedly pineal hyperplasia.

Fasting hypoglycaemia and multiple abnormal features are specific to leprechaunism. All five sibs in this report showed these features (table 1), but the absence of severe growth retardation, long survival, and normal subcutaneous tissue make it difficult to establish a definite diagnosis of classical leprechaunism in this family. However, there are published reports of patients with leprechaunism who had
normal birth weight and long survival, yet other clinical features and molecular studies confirmed the diagnosis of leprechaunism.\textsuperscript{4,15} At the molecular level, the main defect in leprechaunism is a mutation in the insulin receptor gene. Many different mutations in this gene have been described (table 2), but all result in either a decrease in the number of insulin receptors of the target cells\textsuperscript{6} or in a defect in the insulin receptor function itself.\textsuperscript{19,20}

All these patients studied with leprechaunism have had two mutant alleles of the insulin receptor gene. However, some patients were heterozygous for two different mutations and some homozygous for the same mutation,\textsuperscript{15,21,22} but there has been no attempt to correlate phenotype with genotype.

In addition defects in the functions of IGF-1 and EGF receptors have also been described in fibroblasts from some patients,\textsuperscript{24,25} but there is no description of mutations in the genes encoding IGF-1 and EGF receptors.\textsuperscript{20} Kadowaki et al\textsuperscript{20} suggested that the insulin receptor gene may regulate the function of receptors for IGF-1 and EGF so that mutations in the insulin receptor gene might indirectly impair the function of receptors for other growth factors, but why this should be the case in some patients and not others is not clear. There is evidence to suggest that at high concentrations insulin acts through the IGF-1 receptor as a mitogen stimulating thymidine uptake, DNA synthesis, and cell replication. The selective action of insulin on some tissues is related to the density or affinity of IGF-1 receptors.\textsuperscript{6}

The ovary, heart, kidney, and other vascular endothelium have IGF-1 receptors\textsuperscript{26-28} while human fat cells do not.\textsuperscript{29} This could explain the ovarian enlargement, myocardial hypertrophy, and the kidney enlargement reported in some patients with leprechaunism. In addition, Funakoshi et al\textsuperscript{20} reported a case with severe intrauterine growth retardation and abnormalities of IGF-1/SMC receptor functions. They suggested that the severe intrauterine growth retardation was the result of abnormality or immaturity of the IGF-1 receptors.

All the five children in this report have myocardial hypertrophy, kidney enlargement, and ovarian enlargement in the female, but no intrauterine growth retardation. This would confirm that IGF-1 receptors are normal in this form of leprechaunism. Definitive answers should be provided by the molecular characterisation of the mutation in this family and such studies are under way.

The author acknowledges with thanks Dr Simeon Taylor for conducting the molecular studies on this family.

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J Med Genet 1993 30: 470-475
doi: 10.1136/jmg.30.6.470

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Correction

In the paper ‘A syndrome of insulin resistance resembling leprechaunism in five sibs of consanguineous parents’ (J Med Genet 1993;30:470–5), fig 2B is of IV.1 and fig 3B of IV.5.