Respiratory chain complex V deficiency due to a mutation in the assembly gene ATP12

L De Meirleir, S Seneca, W Lissens, I De Clercq, F Eyskens, E Gerlo, J Smet, R Van Coster

In patients with mitochondrial encephalomyopathies an increasing number of causative gene defects have been detected. The number of identified pathogenic mitochondrial DNA mutations has largely increased over the past 15 years. Recently, much attention has turned to the investigation of nuclear oxidative phosphorylation (OXPHOS) gene defects. Within the OXPHOS defects, complex V deficiency is rarely found and, so far, these defects have only been attributed to mutations in the mitochondrial MTATP6 gene.

Mutation analysis of the complete coding regions at the cDNA level of the nuclear ATP11, ATP12, ATP6, ATP8, and ATP13 genes and the mitochondrial MTATP6 and MTAT8 genes was undertaken in two unrelated patients. Blue Native polyacrylamide gel electrophoresis followed by catalytic staining had already documented their complex V decreased activity.

Extensive molecular analysis of five nuclear and two mitochondrial genes revealed a mutation in the ATP12 assembly gene in one patient. This mutation is believed to be the cause of the impaired complex V activity. To our knowledge, this is the first report of a pathogenic mutation in a human nuclear encoded ATPase assembly gene.

The respiratory chain/oxidative phosphorylation system consists of five multisubunit complexes, four of which, complexes I, II, III, and IV, cooperate to generate a proton gradient across the mitochondrial inner membrane. Complex V (ATP synthase or ATPase) couples proton flow from the intermembrane space back to the matrix by the conversion of ADP and inorganic phosphate to ATP. ATP synthase comprises an integral membrane component F0 and a peripheral moiety F1. All five subunits of F1 (α, β, γ, δ, ε) and most F0 subunits of the mammalian ATP synthase are nuclear encoded. Only two F0 proteins (ATP6 and 8) are encoded by mitochondrial DNA. Besides the nuclear genes coding for the subunits of the ATPase complex, several assembly genes have been studied in yeast and their human analogues have been described. Complete loss of the ATP synthase enzyme activity is probably not compatible with life. However, partial loss reflected by a lower amount of functional ATP synthase has been associated with human disease. This was the case for mutations in MTATP6, a mitochondrial encoded component, which causes ATP synthase deficiency. No pathogenic mutations involving nuclear encoded subunits have yet been found. We studied two unrelated children with decreased complex V activity and report on a mutation in the ATP12 assembly gene in one patient.

PATIENTS

Patient 1 was a full-term baby girl, born with a birth weight of 2.550 kg and head circumference of 30.5 cm, the first child of healthy consanguineous parents of Moroccan origin. Family history was unremarkable. The girl presented with dysmorphic features, including a large mouth, prominent nasal bridge, micrognathia, rocker bottom feet, and flexion contractures of the limbs associated with camptodactyly. She was hypertonic and did not suck well. The liver was enlarged. Initial investigation including karyotype and skeletal x ray was normal. The kidneys were hypoplastic. Clinically, the presentation resembled COFS (cerebro-oculofacioskeletal) syndrome (Pena-Shokeir type II), except for the absence of microphthalmia and cataracts. Metabolic screening showed increased urinary lactate, fumarate, methylglutaconic acid, and amino acids. In CSF, lactate was 2.9 mmol/l (normal <2), pyruvate was normal. In plasma, amino acids were normal but lactate fluctuated between 2.7 and 10 mmol/l (normal <2.2) and pyruvate between 0.08 and 0.12 mmol/l (normal 0.03–0.10).

Cerebral magnetic resonance imaging revealed marked cortical–subcortical atrophy, dysgenesis of the corpus callosum with absent anterior genu and rostrum, and hypoplasia of white matter. Within months, the basal ganglia and thalami became atrophic. At the age of 6 months, a muscle and liver biopsy was performed and frozen immediately at −80°C. A skin specimen was used for fibroblast culture. The child had a severe developmental delay with seizures and failure to thrive, and died at the age of 14 months from intercurrent infection.

Patient 2 was born at 37 weeks of gestational age from non-consanguineous white parents. A selective caesarian section was performed because of breech presentation. Birth weight was 2.300 kg, height 48 cm and head circumference 32.7 cm. Apgar scores were 6, 8 and 8, after 1, 5, and 10 minutes respectively. Within the first day the patient started grunting and required supplementary oxygen. The liver was enlarged. On day 2, a sudden deterioration occurred, with desaturation and poor peripheral circulation necessitating endotracheal intubation, artificial ventilation, and administration of a plasma expander. Cerebral ultrasound showed intraventricular haemorrhage and periventricular hyperechogenic zones. The patient started having convulsions and was treated by intravenous administration of phenobarbital. His blood lactate was 50 mmol/l (normal <2). A cardiac arrest necessitated administration of adrenaline and heart massage. Cardiac activity resumed after infusion of sodium bicarbonate. The patient remained hypotensive, had poor peripheral circulation, and dopamine was administered continuously. Ultrasound examination of the heart revealed a small left ventricle. Severe metabolic acidosis persisted despite all efforts to correct it. Following...

Abbreviations: BN-PAGE, Blue Native polyacrylamide gel electrophoresis; CRM, cross-reacting material; OXPHOS, oxidative phosphorylation
another episode of cardiac arrest the patient died on day 3. Immediately post-mortem an autopsy was performed. Skeletal muscle (quadriceps), liver, and tissue specimen were taken and frozen in liquid nitrogen. Microscopic studies were not performed. A skin specimen was used for fibroblast culture.

MATERIALS AND METHODS
Spectrophotometric assays were performed as previously described.

Isolation of mitochondria, solubilisation of oxidative phosphorylation complexes, SDS-PAGE, BN-PAGE and catalytic staining were performed as reported. Oxidative phosphorylation complexes, SDS-PAGE, BN-15% polyacrylamide gel.

detected by digestion of the PCR fragment and analysis on a 9
(5
change in exon 3 of the primers were used. (PCR sequences and PCR conditions are described. Isolation of mitochondria, solubilisation of the complexes separated by BN-PAGE using specific antibodies against several complex V subunits, whereas CRM for complex III was normal (fig 1A). A significant decrease of several individual subunits of complex V was also documented (fig 2B). Two dimensional electrophoresis (BN-PAGE/SDS-PAGE) showed that all major components of complex V (α, β, γ, b, OSCP, d) could not be visualised in heart muscle from the patient (data not shown). Molecular analysis for mitochondrial DNA mutations in the MTP6 and MTP8 subunits was negative. Screening genes α, β, and γ of F1, F2, and assembly genes ATP11 and ATP12 did not reveal any mutation.

DISCUSSION
Mutations have been described in structural subunits of complex I and complex II. Mutations in these genes are responsible for complex IV disrupted activity, while mutations in the BCS1L gene cause complex III failure in patients.

Interestingly, Holme described in 1992 the association of methylglutaconic aciduria and ATP synthase deficiency. The child presented with severe lactic acidosis, hypertrophic cardiomyopathy, and severe mitochondrial ATP synthase deficiency. Methylglutaconic aciduria seems to be a marker of inner mitochondrial membrane dysfunction and can also be seen in other OXPHOS defects. Complex V deficiency is more difficult to detect using spectrophotometric methods but easier with BN-PAGE followed by catalytic staining.

F₂F₃-ATP synthase is comprised of at least fourteen nuclear-encoded subunits and two mitochondrial DNA

Table 1 Activity levels of OXPHOS enzymes in the patients’ samples

<table>
<thead>
<tr>
<th>Tissue</th>
<th>Complex I</th>
<th>Complex II</th>
<th>Complex II-III</th>
<th>Complex IV</th>
<th>Complex V</th>
<th>Citrate synthase</th>
</tr>
</thead>
<tbody>
<tr>
<td>Skeletal muscle</td>
<td>134</td>
<td>131</td>
<td>139</td>
<td>541</td>
<td>119</td>
<td>1011</td>
</tr>
<tr>
<td>Liver (mitochondria)</td>
<td>56</td>
<td>225</td>
<td>200</td>
<td>486</td>
<td>42</td>
<td>471</td>
</tr>
<tr>
<td>Liver (mitochondria)</td>
<td>113</td>
<td>121</td>
<td>192</td>
<td>939</td>
<td>209</td>
<td>1279</td>
</tr>
<tr>
<td>Liver (mitochondria)</td>
<td>194</td>
<td>190</td>
<td>364</td>
<td>87</td>
<td>223</td>
<td></td>
</tr>
</tbody>
</table>

Specific activities are expressed as nmol of substrate/min/mg protein, control values as median (percentile 5–95).
Figure 1  Catalytic staining following separation of the OXPHOS complexes by BN-PAGE showing severely decreased intensities of the bands corresponding to complex V in patients 1 and 2. In skeletal muscle from patient 1, the intensity of staining for complex V is relatively higher than in liver and in cultured skin fibroblasts. The bands corresponding to complexes I, II, and IV in both patients are comparable to the control.

Figure 2  (A) Immunoblotting of complexes III and V following separation by BN-PAGE. Panel 1: a mixture of specific antibodies against the α subunit of complex V and against the core 2 subunit of complex III were used. CRM against complex V was undetectably low in the liver of patient 1 and in heart and skeletal muscle of patient 2, whereas CRM against complex III in both patients was comparable with the control. Panel 2: antibodies were stripped off and the membranes were blotted again using specific antibodies against the β subunit of complex V. Complex V could not be visualised in the liver of patient 1 or in the heart or skeletal muscle of patient 2. Panel 3: membranes were re-used and blotted with specific antibodies against the d subunit of complex V. Complex V could not be visualised in heart or skeletal muscle of patient 2. (B) Immunoblot visualizing subunits of complex V in liver and heart muscle. Mitochondrial proteins were separated using SDS-PAGE. A significant decrease of CRM for the α, β, and OSCP subunits of complex V is shown in patient 1 and patient 2, compared with the control. CRM for the 37 kDa subunit of complex I in both patients was comparable with the control, as was CRM for the SDHb (Fp) subunit of complex II and subunit 3 of complex IV.
Clinical syndrome manifests as NARP (neuropathy, ataxia, retinitis pigmentosa) or the more severe MILS (maternally inherited Leigh's syndrome). Most frequently, these two related mitochondrial disorders are due to a T8993G or T8993C point mutation. In one patient with complex V deficiency presenting in the neonatal period with congenital hyperlactacidaemia, Housteck et al showed, using complementation studies, that the defect was located in the nuclear DNA. Electrophoresis and western blot analysis revealed a reduction of complex V whereas the other OXPHOS complexes were present at normal levels. Inspired by the major involvement of assembly genes in complex IV deficiency and failing to find any mutations in the genes \(\alpha, \beta,\) and \(\gamma\), we initiated a mutation analysis of the assembly genes \(ATP11\) and \(ATP12\) in two patients for whom the BN-PAGE gel had shown a severely decreased activity, an indicator of catalytic deficiency.

Like many other mitochondrial proteins, the \(F_1\) subunits are imported into the matrix compartment as unfolded protein chains. Heat shock proteins HSP60 and HSP10 facilitate their folding. In *Saccharomyces cerevisiae*, the final steps in the formation of a fully functional \(F_1\) require two proteins called Atp11p and Atp12p. It has been shown that they have an essential and indispensable role in the assembly of the \(F_1\) moiety of the ATP synthase. Yeast mutants that are deficient for either Atp11p or Atp12p accumulate both the \(F_1\) and \(\beta\) subunits in large protein aggregates inside mitochondria instead of forming the enzyme oligomer. These strains are respiratory deficient and fail to grow on non-fermentable carbon sources. Interestingly, in addition to being ATPase deficient, the mutant strains had lowered activities of cytochrome c oxidase and coenzyme QH\(_2\)-cytochrome c reductase. This was also found in our patient. ATP11p was shown to bind the \(\beta\) subunit of \(F_1\), while ATP12p interacts specifically with the \(F_1\) subunit. These studies provide the basis for a model of \(F_1\) assembly in which Atp12p is released from the \(\alpha\) subunit in exchange for the \(\beta\) subunit to form the interhelicase that contains the non-catalytic adenine nucleotide binding site. This binding mechanism is thought to protect the \(F_1\) subunits from forming non-productive large complexes during the assembly of the enzyme oligomer.

The human \(ATP11\) and \(ATP12\) genes were only recently identified by Wang et al. The gene products of the human cDNAs act in a manner analogous to their yeast counterparts. Moreover, human \(ATP12\) cDNA complements a yeast \(ATP12\) disruption mutant. Both the \(ATP11\) and \(ATP12\) genes are broadly conserved in eukaryotes and are expressed in a wide range of tissues. The full length cDNA has been characterised and the gene assigned to chromosome 17p11.2.

The ATP11 and ATP12 assembly factors are not members of the major protein chaperone families (for example HSP60), which work for different substrates. Instead, the action of Atp11p and 12p is limited to the biogenesis pathway of the ATP synthase. Neither Atp11p nor Atp12p has a significant homology with other proteins. Elaborated studies of Ackerman and co-workers have already revealed sequence elements in Atp12p that are functionally relevant. Deletion analysis disclosed that some regions could be removed from the mature protein without completely losing the action of \(F_1\) assembly in vivo. The functional domain for Atp12p maps between Asn181 and Val306 in the carboxyl half of the protein. The terminal 18 amino acids of the protein are involved in oligomerisation with other proteins. The precise role of the amino acids in the domain enclosed by the mitochondrial targeting signal on the left and the functional domain on the right is not clear. Truncated Atp12p proteins, with as many as 180 residues of the amino terminal boundary deleted, but with an intact targeting signal, are still capable of conferring some respiratory activity to the yeast cell. However, it has also been shown that these mutants are temperature sensitive, as the yeast strain is

![Figure 3](http://jmg.bmj.com/)
completely respiratory deficient at 30°C but displays partial respiration function at 23°C. Missense mutations located in this region in the yeast protein are, unfortunately, not available. The mutation found in our patient at position 94 lies adjacent to a strongly conserved glutamine conserved in six studied species (fig 3B). Probably, the Atp12p activity is severely compromised when a neutral polar amino acid (tryptophan) is changed for a basic one (arginine), and is no longer able to mediate proper F1 assembly.

In conclusion, complex V deficiency seems to be an early presenting disease in which lactic acidosis, dysmorphic features, and methyl glutacetic aciduria can be major clues in the diagnosis. To our knowledge, this is the first report of complex V deficiency with a mutation in a nuclear encoded assembly gene for ATPase. It is clear that the search for genes causing respiratory chain disorders has only just begun, and many more mutations will be identified in the near future.

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Authors’ affiliations
L De Meirleir, Department of Paediatric Neurology, University Hospital Vrije Universiteit Brussel (AZK-VUB), Brussels, Belgium
S Seneca, W Lissens, I De Clercq, Department of Medical Genetics, University Hospital, Vrije Universiteit Brussel (VUB), Brussels, Belgium
F Eyksen, Department of Pediatrics and Metabolic diseases, University Hospital, Antwerp, Belgium
E Gerlo, Department of Clinical Chemistry, University Hospital, Vrije Universiteit Brussel (VUB), Brussels, Belgium
J Smet, R Van Coster, Department of Pediatrics, Division of Pediatric Neurology and Metabolism, Ghent University Hospital, Ghent, Belgium

Correspondence to: Dr L De Meirleir, Pediatric Neurology, AZK-VUB, Laarbeeklaan 101, B-1090 Brussels, Belgium; linda.demeirleir@az.vub.ac.be

REFERENCES
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