LIPOAMIDE DEHYDROGENASE DEFICIENCY: A NEW CAUSE FOR RECURRENT MYOglobinuria

O.N. ELPELEG, MD, A.B. SAADA, PhD, A. SHAAG, MSc, J.Z. GLUSTEIN, MD, W. RUITENBEEK, PhD, I. TEIN, MD, and J. HALEVY, MD

Inborn errors of pyruvate metabolism and the mitochondrial respiratory chain are rarely associated with recurrent myoglobinuria. A defect in the proteins containing iron-sulfur clusters, including components of the Krebs cycle and the mitochondrial respiratory chain, was found in 1 patient and deletions of mitochondrial DNA (mtDNA) were reported in 3 patients. Severe coenzyme Q deficiency in muscle was detected in 2 sisters who suffered since early childhood from recurrent myoglobinuria and myopathy, short stature, seizure disorder, and cerebellar symptoms. We describe a 37-year-old man with recurrent myoglobinuria and lactic acidemia due to deficiency of the mitochondrial enzyme lipoamide dehydrogenase (LAD).

CASE REPORT

A 37-year-old man of Ashkenazi-Jewish origin developed generalized weakness, recurrent vomiting, and weight loss of 10 kg over a period of 2 months. His parents, 2 sisters, and 4 children were healthy. He suffered several episodes of muscle weakness associated with "anicteric hepatitis" in the past 17 years. Prior to admission, he had elevated alanine aminotransferase (ALT) (peak value, 556 U/L; control, 5-50 U/L) and aspartate aminotransferase (AST) (231 U/L; control, 7-56 U/L). On admission, he had symmetric generalized hyporeflexia. Electromyography revealed low-voltage polyphasic motor unit potentials. Creatine kinase (CK) (11277 U/L; control, 30-150 U/L), AST (400 U/L), and ALT (757 U/L) were elevated. After administration of chlorpromazine (12.5 mg IV) to alleviate vomiting, he was found pale, tachypneic, and confused, with severe metabolic acidosis (blood pH 6.8, pCO2 14). Serum lactate was 29.7 mmol/L (control, 0.9-2.1 mmol/L), β-hydroxybutyrate 2.77 mmol/L (control, 0.05-0.2 mmol/L), acetoacetate 0.28 mmol/L (control, 0.02-0.15 mmol/L), CK 71000 U/L, serum carnitine 167 µmol/L (control, 30-60 µmol/L), AST 591 U/L, and ALT 1640 with prolongation of the prothrombin time. The branched-chain amino acids in plasma were 1.5-2.5 times the upper limit of the control range. In the urine, myoglobin was strongly positive and organic acid analysis revealed massive excretion of lactic and β-hydroxybutyric acids, low-normal levels of Krebs cycle components, and no evidence of fatty acid oxidation defect. A biopsy of the quadriceps revealed small cytoplasmic lipid droplets. Total muscle carnitine was 3.3 µmol/g tissue (control, 3.1 µmol/g tissue) and nonesterified carnitine was 3.1 µmol/g tissue (control, 2.2-4.2). The patient recovered with complete normalization of all biochemical abnormalities, but reported occasional vomiting and mild exertional myalgia. Therapy with lipoic acid was recommended but compliance was poor. He was rehospitalized 5 months later because of vomiting, weakness, and elevation of lactate (24.2 mol/L), ammonia (150 µmol/L), and CK (1129 U/L). An intractable metabolic acidosis developed and the patient expired 12 h after admission. Consent for autopsy was not given.

METHODS

In the mitochondrial enriched fraction of the muscle, the activity of citrate synthase, reduced form of nicotinamide adenine dinucleotide (NADH)-
ferricyanide reductase, antimycin-sensitive succinate-
co
cytochrome c oxidoreductase, succinate-
St
ubiquinone reductase, ubiquinol-cytochrome c
reductase, pyruvate dehydrogenase complex (PDHc),
and cytochrome c oxidase were determined. The ac-
tivity of carnitine palmitoyltransferase was measured
in the 600-g supernatant of the muscle and the ac-
tivities of PDHc, pyruvate decarboxylase, LAD, and
\( \alpha \)-ketoglutarate dehydrogenase complex (\( \alpha \)KGDHc)
were determined in muscle homogenate and in fi-
broblasts.\textsuperscript{5,14,16,19}

PDHc antiserum was produced in rabbits as de-
scribed by De Marcucci et al.\textsuperscript{1} LAD antiserum was
produced by immunization of rabbits with 0.5 mg of
LAD (Sigma Chemicals, St. Louis, MO) in complete
Freund's adjuvant, followed by three boosts in in-
complete Freund's adjuvant. Muscle mitochondrial
fractions of the patient and controls were separated
by sodium dodecyl sulfate/polyacrylamide gel elec-
trophoresis (SDS-PAGE), transferred to polyvinyl-
idenefluoride (PVDF) membrane, and incubated
with anti-PDHc or anti-LAD antisera. Antigen-
\footnotemark[1]
antibody complexes were visualized using the alkali-
ne-phosphatase conjugate substrate kit (Bio-Rad
Laboratories, Richmond, CA).

Genomic DNA was isolated and cDNA was pro-
duced from fibroblasts of the patient and 5 controls.
Direct sequencing of the entire coding region of the
LAD gene and of a genomic DNA fragment up to
nt -207 was performed as previously described.\textsuperscript{3}

**RESULTS**

The enzymatic analysis revealed normal or elevated
activities of all mitochondrial enzymes in muscle tis-
tue with the exception of PDHc (11% of the control
mean), \( \alpha \)KGDHc (5%), and LAD (21%). The activity
of LAD in fibroblasts was also reduced (11.5 nmol/
min/mg protein; control, 83 ± 7.6).

The western blot disclosed reduced amount of the
LAD protein (25% of the control) and normal
amounts of pyruvate decarboxylase subunits (\( E_\text{1a} \) and
\( E_\text{1b} \)), transacetylase (\( E_\text{2} \)), and the X protein (Fig. 1).
No mutations were detected in the coding region
of the cDNA of LAD and the 5' flanking region up to
nt -207.

**DISCUSSION**

Recurrent myoglobinuria was reported to accom-
pany at least 17 inborn errors of metabolism. These
include six heritable defects in glycogenolysis or gly-
colysis, six in the pathway of short-, medium-, or
long-chain fatty acid oxidation, one in the pentose
phosphate pathway, one in the purine nucleotide
cycle, and three in the respiratory chain, Krebs cycle,
and mtDNA. Partial deficiency of carnitine palmito-
yltransferase II is the most common cause of recur-
rent rhabdomyolysis.\textsuperscript{2,12,13,16,20}

We attribute the recurrent myoglobinuria and
lactic acidemia to LAD deficiency in this patient.
LAD is one of the three catalytic proteins of the

\begin{figure}
\centering
\includegraphics[width=\textwidth]{figure1}
\caption{Western blot analysis: Lanes 1 and 4: 5 \( \mu \)g of mitochondrial protein of the patient; lanes 2, 3, 5, and 6: 5 \( \mu \)g of mitochondrial protein of 2 controls with citrate synthase activity which is similar to that of the patient; lane 7: procine LAD. Lanes 1–3 were exposed to
anti-PDHc antiserum (this serum does not react with LAD). Lanes 4–7 were exposed to anti-LAD antiserum.}
\end{figure}
PHDc which converts pyruvate to acetyl-CoA and a component of two other α-keto acid dehydrogenase complexes, αKGDHc and branched-chain α-keto acid dehydrogenase complex. Thus, the lactic acidemia, elevation of branched-chain amino acids, and reduction of both PDHc and αKGDHc as seen in our patient can all result from a single defect in LAD.

Because abnormalities of mitochondrial enzymes are not commonly associated with rhabdomyolysis, we considered the possibility that the deficiency of dehydrogenase complex. Thus, the lactic acidemia, which was also present in fibroblasts which were propagated in culture, excluding a transient inhibition of the enzyme activity.

The few patients with LAD deficiency previously reported had a clinical picture which was in no way similar to that of our patient. Their disease was characterized by progressive neurological deterioration and lactic acidemia, and most of them died within the first few years of life.1,9,10,14,15,17 Peripheral muscle involvement was not previously documented. We have diagnosed several children with LAD deficiency whose presentation in infancy was of hepatocellular failure, sparing central nervous system and muscle.3,4 In some of them, sequencing of the LAD gene revealed an insertion mutation within the leader sequence. In the present patient, no mutation was identified in the entire coding region of the LAD gene or in the immediate 5'-flanking region. In view of the reduced quantity of the enzymatic protein in the muscle tissue, the disease-causing mutation may interfere with the translation or transcription process; such a mutation may reside in fragments -208 to -583 and -646 to -864, which were shown to contain positive transcriptional regulatory elements and were not yet analyzed by us.7

In summary, the present report widens the clinical spectrum of LAD deficiency to include adult-onset recurrent myoglobinuria, and indicates that LAD deficiency should be included in the differential diagnosis.

REFERENCES


8. Keightley JA, Hoffbuhler KC, Burton MD, Salas VM, Johnston JMF, Guggenbichler JP, Janssen AJM, Bakkeren JAJM: Deficiency of the human dihydrolipoamide dehydrogenase gene or in the immediate 5'-flanking region. In view of the reduced quantity of the enzymatic protein in the muscle tissue, the disease-causing mutation may interfere with the translation or transcription process; such a mutation may reside in fragments -208 to -583 and -646 to -864, which were shown to contain positive transcriptional regulatory elements and were not yet analyzed by us.7


