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# Pathogenic variants in *SQOR* encoding sulfide:quinone oxidoreductase are a potentially treatable cause of Leigh disease

Marisa W. Friederich<sup>1,2</sup> | Abdallah F. Elias<sup>3</sup> | Alice Kuster<sup>4,5</sup> |  
 Lucia Laugwitz<sup>6</sup> | Austin A. Larson<sup>1</sup> | Aaron P. Landry<sup>7</sup> |  
 Logan Ellwood-Digel<sup>1</sup> | David M. Mirsky<sup>8</sup> | David Dimmock<sup>9</sup> | Jaclyn Haven<sup>3</sup> |  
 Hua Jiang<sup>1</sup> | Kenneth N. MacLean<sup>1</sup> | Katie Styren<sup>3</sup> | Jonathan Schoof<sup>3</sup> |  
 Louise Goujon<sup>4,10</sup> | Thomas Lefrancois<sup>11</sup> | Maïke Friederich<sup>1</sup> |  
 Curtis R. Coughlin II<sup>1</sup> | Ruma Banerjee<sup>7</sup> | Tobias B. Haack<sup>5,12</sup> |  
 Johan L. K. Van Hove<sup>1,2</sup>

<sup>1</sup>Section of Clinical Genetics and Metabolism, Department of Pediatrics, University of Colorado, Aurora, Colorado

<sup>2</sup>Department of Pathology and Laboratory Medicine, Children's Hospital Colorado, Aurora, Colorado

<sup>3</sup>Department of Medical Genetics, Shodair Children's Hospital, Helena, Montana

<sup>4</sup>Department of Neurometabolism, University Hospital of Nantes, Nantes, France

<sup>5</sup>INRAE, UMR1280, PhAN, Nantes Université, Nantes, France

<sup>6</sup>Institut für Medizinische Genetik und Angewandte Genomik, Universitätsklinikum, University of Tübingen, Tübingen, Germany

<sup>7</sup>Department of Biological Chemistry, University of Michigan, Ann Arbor, Michigan

<sup>8</sup>Department of Radiology, University of Colorado, and Children's Hospital Colorado, Aurora, Colorado

<sup>9</sup>Rady Children's Institute for Genomic Medicine, San Diego, California

<sup>10</sup>Service de Génétique Clinique, University Hospital of Rennes, Rennes, France

<sup>11</sup>Pediatric Radiology, University Hospital of Nantes, Nantes, France

<sup>12</sup>Centre for Rare Diseases, University of Tübingen, Tübingen, Germany

## Correspondence

Johan L. K. Van Hove, Department of Pediatrics, Section of Clinic Genetics and Metabolism, University of Colorado, Education 2 South, L28-4114, 13121 East 17th Avenue, Aurora, CO 80045.  
 Email: johan.vanhove@childrenscolorado.org

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## Abstract

Hydrogen sulfide, a signaling molecule formed mainly from cysteine, is catabolized by sulfide:quinone oxidoreductase (gene *SQOR*). Toxic hydrogen sulfide exposure inhibits complex IV. We describe children of two families with pathogenic variants in *SQOR*. Exome sequencing identified variants; *SQOR* enzyme activity was measured spectrophotometrically, protein levels evaluated by western blotting, and mitochondrial function was assayed. In family A, following a brief illness, a 4-year-old girl presented comatose with lactic acidosis and multiorgan failure. After stabilization, she remained comatose, hypotonic, had neurostorming episodes, elevated lactate, and Leigh-like lesions on brain

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imaging. She died shortly after. Her 8-year-old sister presented with a rapidly fatal episode of coma with lactic acidosis, and lesions in the basal ganglia and left cortex. Muscle and liver tissue had isolated decreased complex IV activity, but normal complex IV protein levels and complex formation. Both patients were homozygous for c.637G > A, which we identified as a founder mutation in the Lehrerleut Hutterite with a carrier frequency of 1 in 13. The resulting p. Glu213Lys change disrupts hydrogen bonding with neighboring residues, resulting in severely reduced SQOR protein and enzyme activity, whereas sulfide generating enzyme levels were unchanged. In family B, a boy had episodes of encephalopathy and basal ganglia lesions. He was homozygous for c.446delT and had severely reduced fibroblast SQOR enzyme activity and protein levels. SQOR dysfunction can result in hydrogen sulfide accumulation, which, consistent with its known toxicity, inhibits complex IV resulting in energy failure. In conclusion, SQOR deficiency represents a new, potentially treatable, cause of Leigh disease.

#### KEYWORDS

complex IV, hydrogen sulfide, Leigh disease, sulfide:quinone oxidoreductase, treatment

## 1 | INTRODUCTION

Despite the purported role of hydrogen sulfide (H<sub>2</sub>S) as a signaling molecule with multiple physiologic functions, few inherited disorders have been described involving its metabolism or function.<sup>1,2</sup> Here, we describe a new disorder of H<sub>2</sub>S metabolism.

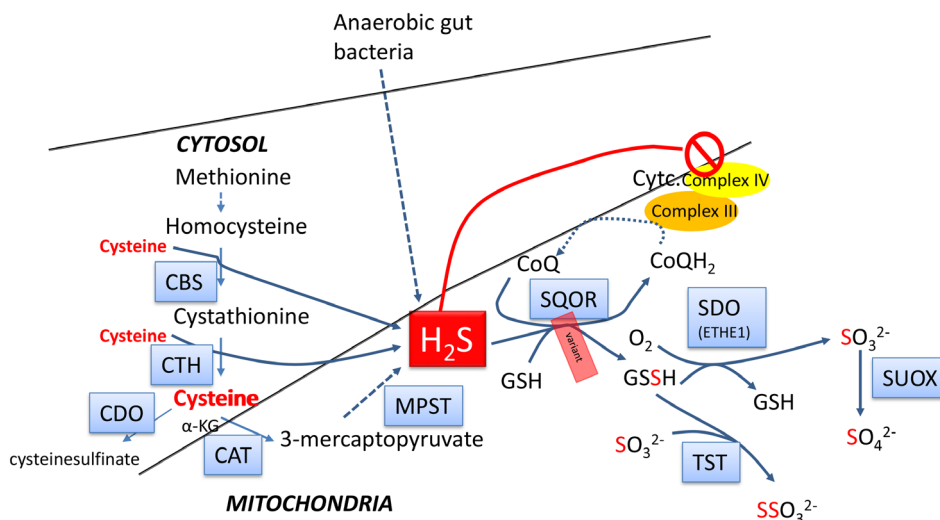
In brain, H<sub>2</sub>S increases cAMP, stimulates NMDA-receptor mediated responses including long-term potentiation, and stimulates calcium influx in astrocytes. In the vascular system, it promotes vasodilatation and angiogenesis.<sup>1,2</sup> It upregulates antioxidant systems, anti-inflammatory, and cytoprotective genes.

The toxicity of excess H<sub>2</sub>S, such as occurs during exposure in industrial accidents, results in part from inhibiting cytochrome c oxidase, including by coordination with the iron of heme a.<sup>3</sup> It changes the ventilatory drive possibly by inhibiting carbonic anhydrase, and can cause pulmonary edema and cardiac depression. Chronic H<sub>2</sub>S accumulation has been implicated in ethylmalonic encephalopathy and related to inhibition of short chain acyl-CoA dehydrogenase and loss of cytochrome c oxidase.<sup>4-7</sup> Finally, H<sub>2</sub>S has been implicated in the pathophysiology of primary coenzyme Q deficiency.<sup>8,9</sup>

The metabolism of H<sub>2</sub>S is complex (Figure 1). At physiological pH, H<sub>2</sub>S consists of dissolved gaseous H<sub>2</sub>S in equilibrium with HS<sup>-</sup>. It is primarily generated from

cysteine via the activity of three enzymes. In most tissues, the majority of H<sub>2</sub>S production is derived via α,β-elimination by cystathionine γ-lyase (CTH) upon L-cysteine to form L-serine and H<sub>2</sub>S.<sup>10,11</sup> Also, H<sub>2</sub>S is formed in the condensation reaction of cystathionine β-synthase (CBS) when L-serine is substituted by L-cysteine. Further, L-cysteine is transaminated by cysteine aminotransferase (CAT) with α-ketoglutarate forming 3-mercaptopyruvate, which is then acted upon by mercaptopyruvate sulfurtransferase (MPST) to release H<sub>2</sub>S in mitochondria and in the cytosol.<sup>12</sup> In the brain, D-cysteine can be oxidized by D-amino-oxidase to mercaptopyruvate, the substrate for MPST.<sup>12</sup> Finally, gut bacteria produce H<sub>2</sub>S, which is rapidly metabolized in the intestinal epithelium.<sup>7,13-15</sup>

H<sub>2</sub>S is primarily oxidized via sulfide:quinone oxidoreductase, an inner mitochondrial membrane protein encoded by the SQOR gene (previously called *SQRDL*) (OMIM# 617658). A cysteine persulfide is transiently formed in the SQOR active site while electrons from H<sub>2</sub>S oxidation reduce coenzyme Q via the cofactor FAD.<sup>16</sup> The sulfane sulfur from this persulfide intermediate is then transferred to the physiological acceptor glutathione forming glutathione persulfide (GSSH).<sup>17,18</sup> In vitro, sulfite, cyanide, and other acceptors can substitute for glutathione forming thiosulfate, thiocyanate, and other products.<sup>17</sup> Glutathione persulfide is oxidized by persulfide dioxygenase (encoded by the gene *ETHE1*,



**FIGURE 1** The metabolism of hydrogen sulfide. Hydrogen sulfide ( $\text{H}_2\text{S}$ ) is formed from cysteine by cystathionine  $\beta$ -synthase (CBS) or by cystathionine- $\gamma$ -lyase (CTH), or after transamination via cysteine aminotransferase (CAT) and the action of mercaptopyruvate sulfur transferase (MPST). Cysteine is also produced by gut bacteria. Cysteine is oxidized by cysteine dioxygenase (CDO) in the pathway to taurine synthesis.  $\text{H}_2\text{S}$  is primarily oxidized by sulfide:quinone oxidoreductase (SQOR) with the electrons transferred to coenzyme Q (CoQ), and with concomitant sulfur transfer to glutathione (GSH) to form glutathione persulfide (GSSH). GSSH is then oxidized to form sulfite by persulfide dioxygenase (SDO, gene *ETHE1*), which is further oxidized to sulfate by sulfite oxidase (SUOX). GSSH can also be converted to thiosulfate by thiosulfate sulfurtransferase (TST). Accumulating hydrogen sulfide can inhibit respiration at the level of complex IV. Other abbreviations used: CoQ is oxidized coenzyme Q, CoQH<sub>2</sub> is reduced coenzyme Q, Cytc is cytochrome c,  $\alpha$ -KG is  $\alpha$ -ketoglutarate,  $\text{SO}_3^{2-}$  is sulfite,  $\text{SO}_4^{2-}$  is sulfate,  $\text{SSO}_3^{2-}$  is thiosulfate. MPST is variably localized in both the mitochondria and cytosol

pathogenic variants in this gene are implicated in ethylmalonic encephalopathy) to sulfite, which can be further oxidized to sulfate by sulfite oxidase or metabolized by thiosulfate sulfurtransferase to thiosulfate.<sup>7</sup>

Leigh disease is a frequent neurologic presentation of genetic defects in mitochondrial bioenergetics.<sup>19</sup> It is highly heterogenous genetically with more than 75 causative genes identified, yet the genetic cause for many patients remains unresolved.<sup>20</sup> Most patients have defects in structural genes of the respiratory chain complex, or in processes involving transcription, translation, or assembly of the protein complexes including their multiple cofactors. Defects in complex IV are a recurring cause of Leigh disease.<sup>21</sup> Very few mitochondrial disorders are related to secondary toxic inhibition of respiratory chain enzyme complexes. Few mitochondrial disorders have an effective treatment option.

Here, we present children from two families who developed Leigh disease. The genetic defect was identified in the *SQOR* gene, and functional assays show abrogated *SQOR* enzyme activity and isolated decreased complex IV activity, but normal complex IV protein levels and complex formation. Recognition of this new genetic cause of Leigh disease is clinically important as it identifies a potentially treatable condition.

## 2 | METHODS

### 2.1 | Subjects

All procedures were done in accordance with the ethical standards of the ethics committee in accordance with the Helsinki Declaration of 1975 and as revised in 2000. Studies at the University of Colorado were done following IRB-approved protocols 07-0386 and 16-0146. Studies at Shodair Children's Hospital in Montana were done according to IRB-approved protocol IRB00008287. For studies of identifiable living human subjects, written informed consent was obtained prior to initiation of research as outlined in the IRB-approved study. The study was approved by the ethics committee of the University of Tuebingen (437/2018BO1). The clinical histories were reviewed by clinical geneticists, and brain magnetic resonance imaging (MRI) studies were reviewed by a pediatric neuroradiologist (D. M.).

### 2.2 | Exome sequencing

Exome sequencing was performed on the index cases and their parents. For an initial estimate of the carrier rate in the Hutterite population, de-identified DNA samples from this population that had been previously submitted

for diagnostic studies for variable indications were screened by high-resolution melt curve analysis and Sanger sequencing as described in Supplemental Methods. The carrier incidence in the general population was identified from the gnomAD database using a methodology as described in Supplemental Methods.<sup>22</sup>

### 2.3 | Analysis of respiratory chain enzyme activities and assembly

Respiratory chain enzyme activities of complexes I, II, II + III, III, IV, and citrate synthase were analyzed in duplicate in post-600× g supernatants isolated from liver and muscle of subject A.II-3 and from fibroblasts of subject B.II-3 as described previously.<sup>23</sup> Ratios of enzyme activities to either citrate synthase or complex II activities were calculated and expressed as Z-scores of the normal distribution established in control samples. Solubilized inner mitochondrial membrane fractions isolated by differential centrifugation from liver and muscle from subject A.II-3 and from fibroblasts from subject B.II-3 were separated by blue native polyacrylamide gel electrophoresis (BN-PAGE) and evaluated by in-gel activity staining assays as described previously.<sup>23</sup>

Complex IV assembly was analyzed in isolated mitochondrial membrane fractions obtained from liver and muscle from subject A.II-3 using nondenaturing clear native PAGE followed by western blot analysis using the antibody against COXIV. Complex I assembly was analyzed in mitochondrial membrane fractions using nondenaturing BN-PAGE followed by western blot analysis using the antibody against NDUFS2, as described previously.<sup>24</sup> In control samples, this assay shows only a holocomplex at 1 MDa and a faint band at 400 kDa in muscle samples, or at 230 kDa in fibroblasts.

### 2.4 | Western blot analysis of mitochondrial proteins including SQOR

In subject A.II-3, complex IV integrity was evaluated using an antibody against COXIV and complex I integrity was evaluated using an antibody against NDUFB8 on western blot analysis. The amount of SQOR protein was determined by western blot analysis on mitochondrial inner membrane fractions. The levels of H<sub>2</sub>S generating enzymes were evaluated by western blot analysis using antibodies to CBS, CTH, MPST, and cysteine dioxygenase (CDO). All methods are described in Supplemental Methods, and the source and specifications of all antibodies used are listed in Supplemental Table S1.

### 2.5 | Analysis of SQOR enzyme activity in liver, blood cells, and fibroblasts

The activity of SQOR was measured spectrophotometrically in mitochondrial membrane fractions isolated from liver and in lysates derived from peripheral blood mononuclear cells (PBMCs) and from fibroblasts as previously described for purified protein with the modifications described in detail in Supplemental Methods.<sup>26</sup> The reaction uses sulfite as acceptor, following coenzyme Q<sub>1</sub> reduction spectrophotometrically for 5 to 8 minutes at 278 nm. The kinetic parameters for human SQOR enzyme were evaluated in human liver and fibroblast samples.

### 2.6 | Protein modeling, purification, and in vitro characterization of the human SQOR mutant

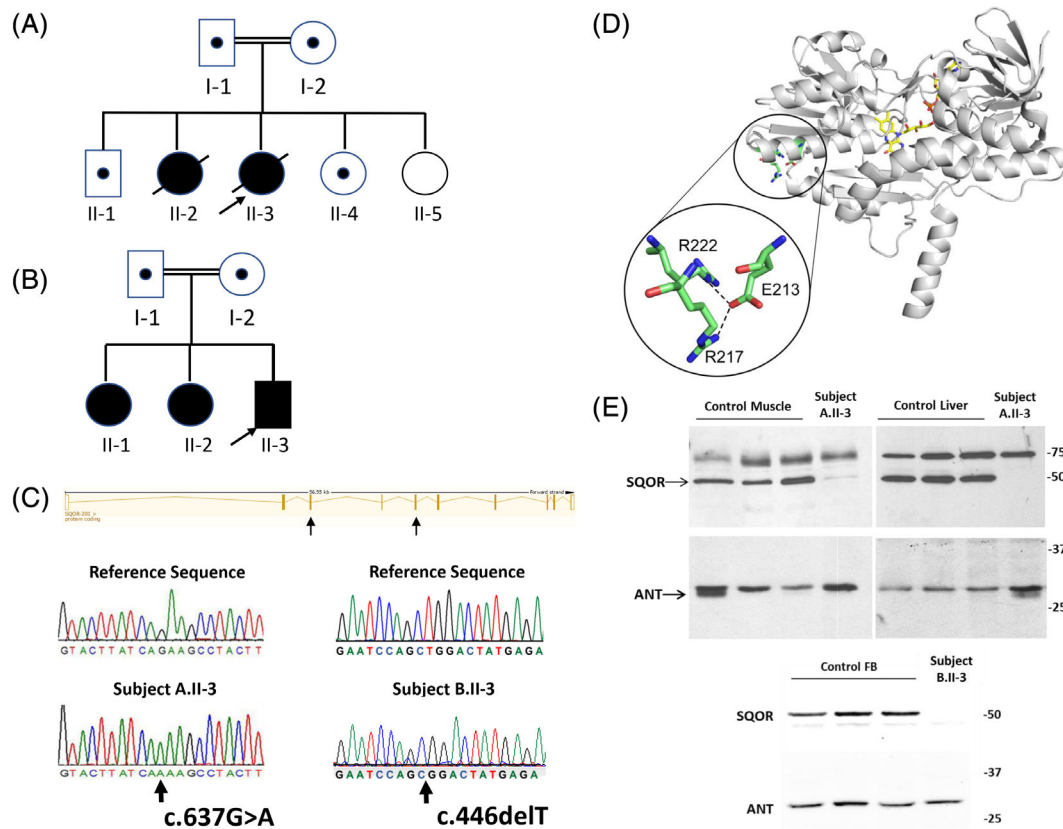
The SQOR variant was mapped on the crystal structure of the human SQOR protein (PDB: 6O1C)<sup>27</sup> using PyMol.<sup>28</sup> The human SQOR mutant protein p.Glu213Lys was purified as reported previously for the wild type enzyme<sup>17</sup> and the absorption spectrum and concentration estimated as described in Supplemental Methods.

## 3 | RESULTS

### 3.1 | Case Reports

#### 3.1.1 | Family A

The proband, subject II-3, was a 4-year-old girl born to consanguineous Lehrerleut Hutterite parents, whose grandparents were first cousins (Figure 2A). She had a history of recovering slower from infections than her siblings, but otherwise had no health concerns including no gastrointestinal symptoms. At onset of the illness, during the day she was more tired and vomited, and in the evening became rapidly encephalopathic with an episode of tonic seizures. Upon presentation, she was comatose with seizure-like movements, and Kussmaul breathing due to lactic acidosis with pH 6.86, anion gap 31, lactate 17.3 mM, and uric acid 7.8 mg/dL. She developed shock, respiratory insufficiency evolving into acute respiratory distress syndrome, and multi-organ failure including liver failure with increased prothrombin time INR 1.76 and decreased clotting factors V and VII, mild hypoglycemia 47 mg/dL, mild hyperammonemia 112 μM, hypoalbuminemia 2.9 g/L, but normal cortisol. With supportive therapy, the multiorgan failure stabilized rapidly over

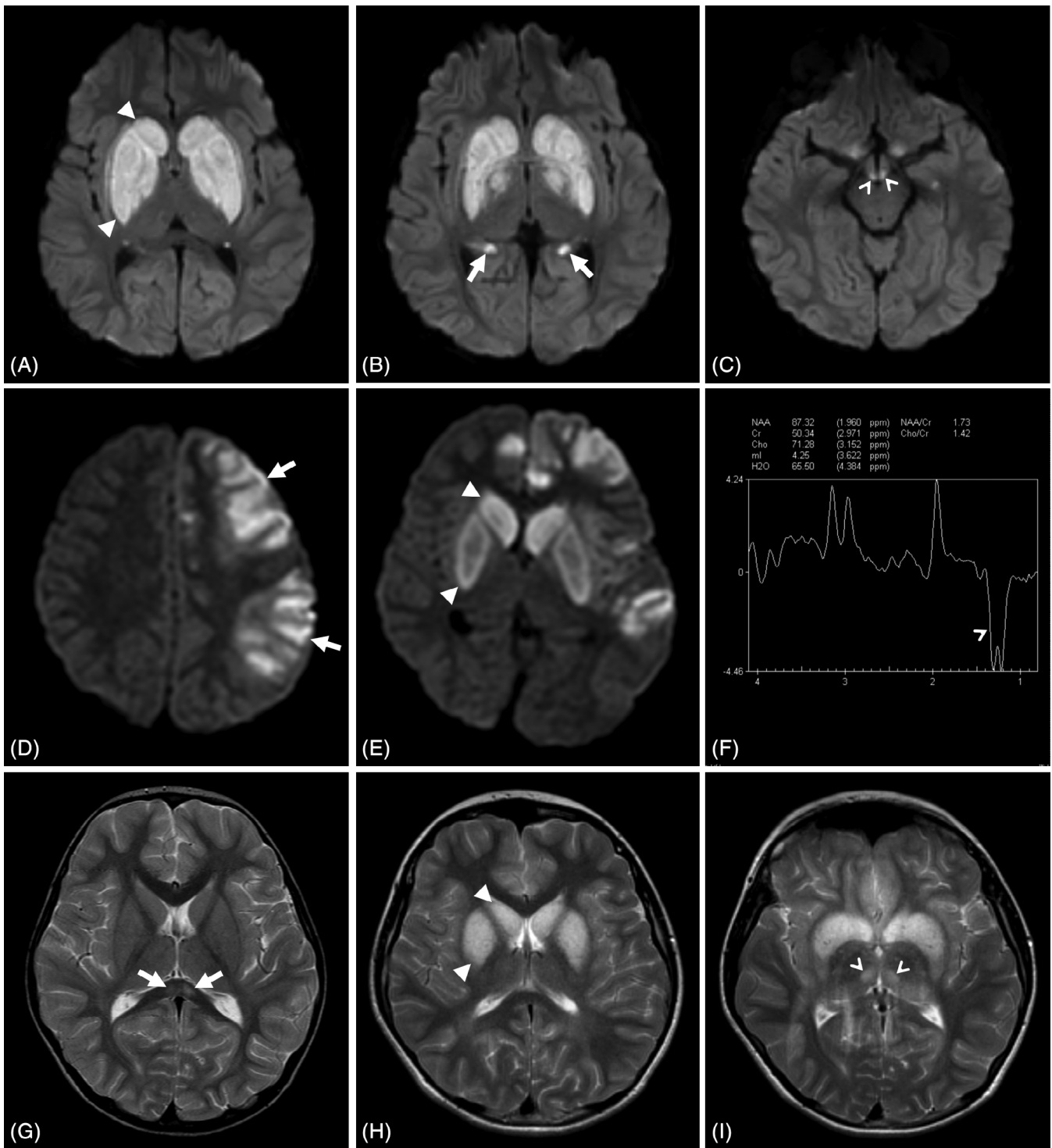


**FIGURE 2** The SQOR pathogenic variant causes lack of protein and enzyme activity. A, In family A, a homozygous pathogenic variant c.637G > A in exon 5 was identified in subjects A.II-2 and A.II-3 indicated by a filled circle, whereas subjects A.I-1, A.I-2, A.II-1, and A.II-4 are carriers for this pathogenic variant. B, In family B, the children are each homozygous for the variant c.446delT whereas the parents are heterozygous carriers. C, Sanger sequencing results of both variants in subjects A.II-3 and B.II-3 are shown. D, The variant c.637G > A codes for p.(Glu213Lys) which is mapped on the crystal structure of the human SQOR protein (PDB: 6O1C). The human SQOR monomer is shown in gray ribbon with the flavin cofactor (yellow) and select residues (green) are shown in stick display. Glu213 is relatively surface exposed. The close-up view shows that the side chain of Glu213 interacts with Arg217 and Arg222. The Glu213Lys mutation is predicted to disrupt electrostatic stabilization of the arginine side chains. E, The SQOR protein (at approx. 50 kDa) is virtually absent in the patient liver and strongly reduced in muscle as shown by western blot, using ANT (approx. 33 kDa) as a loading control in subject A.II-3 and in fibroblast of subject B.II-3

the next days, but coma (Glasgow coma scale [GCS] 5, decorticate posturing) without further seizures, hypoalbuminemia (2.0–2.4 g/dL, normal 3.5–5.2 g/dL) with otherwise normal liver function (transaminases AST 52–88, normal 15–50), and lactic acidosis (3–5 mM with lactate/pyruvate ratio 17 and cerebrospinal fluid [CSF] lactate 3.0 mM) persisted. Creatine kinase peaked at 722 U/L and remained mildly elevated. In subsequent weeks, she had generalized hypotonia, and episodes of neurostorming with tachycardia, hypertension, and potentially pain, partially responding to treatment with phenobarbital, levetiracetam, baclofen, diazepam, and propranolol. She also developed a thrombus in the right iliac to femoral vein requiring anticoagulation. She died a few days after discharge.

Workup for infectious agents and toxins was negative. The acylcarnitine profile, carnitine levels,

3-methylglutaconic acid, thiamin pyrophosphate level, and plasma, urine and CSF amino acids levels were all normal. Urine organic acids showed evidence of lactic acidosis and ketosis. The electroencephalogram showed diffuse slowing and lack of reactivity but no epileptiform activity, and full cardiac evaluation including echocardiogram was normal. Brain MRI on the fourth day of illness showed T2 prolongation and diffusion restriction in the caudate, putamina, and globi pallidi (Figure 3A,B, white arrowheads). There was also diffusion restriction in the posterior part of the hippocampi (Figure 3B white arrows), medial left temporal lobe, posterior inferior thalamus and the mammillary bodies (Figure 3C, white notched arrowheads). Further, there was increased T2 and FLAIR signal in the cerebral peduncles, dentate nuclei, and medial thalami, but brain stem, cerebellum, cortex and white matter were normal. Brain magnetic



**FIGURE 3** Brain magnetic resonance imaging in patients affected by SQOR deficiency. A-C, Axial diffusion-weighted brain MR images of patient A.II-3 demonstrated symmetric restricted diffusion and edema involving the basal ganglia (white arrowheads), hippocampi (white arrows), and mammillary bodies (white notched arrowheads). D-E, Axial diffusion-weighted brain MR images of patient A.II-2 reveal predominantly left-sided restricted diffusion and edema of the cerebral cortices (white arrows) and symmetric restricted diffusion and edema of the basal ganglia (white arrowheads). F, Single voxel proton MR spectroscopy (TE = 144) interrogating the left frontal lobe of patient A.II-2 shows a large inverted lactate doublet at 1.3 ppm. G, In patient B.II-3 at age 4 4/12 years, axial T2-weighted brain MR image illustrated patchy increased signal involving the splenium of the corpus callosum (white arrows). H-I, One year later for the same patient, axial T2-weighted brain MR images show symmetric edema of the basal ganglia (white arrowheads) and anteromedial thalami (white notched arrowheads), all of which had reduced diffusivity (not shown)

resonance spectroscopy (MRS) over the right basal ganglia showed a large lactate peak six times larger than the choline peak with a depressed *N*-acetylaspartate peak. Biopsies of muscle and liver obtained for mitochondrial evaluation had normal histology and electron microscopy including of the mitochondrial architecture, and skin biopsy electron microscopy was normal without evidence of storage products. Nextgen sequencing of mitochondrial DNA showed normal sequence without evidence for deletions.<sup>29</sup> Clinical exome sequencing was reported as negative without pathogenic variants identified in known disease-causing genes.

Subject II-2, the older sister, was generally healthy, but developed migraines 5 months prior to presentation. At age 8 years, she developed a fever, nausea and vomiting, and rapidly became comatose with apnea requiring intubation. She had a large anion gap metabolic acidosis with elevated lactate (peak 22 mmol/L) with lactate/pyruvate ratio 51, with otherwise normal blood values. She was tachycardic but otherwise had normal cardiovascular function. She had depressed encephalopathic function on EEG with spike waves over the left frontal area. She died 3 days later.

On brain MRI, she had restricted diffusion of the left frontoparietal cortex (Figure 3D,E, white arrows) and right frontal cortex (not labeled). She had restricted diffusion of caudate nuclei and putamina (Figure 3E). MRS of the right basal ganglia revealed a large lactate inverted doublet, increased choline and reduced *N*-acetylaspartate signal (Figure 3F, white notched arrowhead).

### 3.1.2 | Family B

Subject II-3 is a male child born to consanguineous parents of Turkish descent (cousins three times removed), who has two healthy sisters (Figure 2B). At age 4 4/12 years, following gastroenteritis with diarrhea and 48 hours fasting, he presented encephalopathic with decreased consciousness (GCS 12), dyskinesia and mydriasis. On admission, he had mild hypoglycemia 50 mg/dL, mild hyperammonemia 100  $\mu$ M, and metabolic acidosis (pH 7.23, bicarbonate 12 mmol/L, base excess  $-14$  mmol/L) with elevated lactate 6.0 mmol/L, and large ketones in urine. He had elevated creatine phosphokinase, myoglobin, and transaminases. Urine organic acids showed elevated lactate, ketones, and dicarboxylic acids. Serum acylcarnitine profile was consistent with ketosis, and plasma amino acids showed elevations of glutamine, lysine, proline, alanine, methionine, and tyrosine with low citrulline, whereas CSF amino acids were generally normal (Supplemental Table S2). Brain MRI showed lesions of the splenium of the corpus

callosum (Figure 3G, white arrows), felt to be clinically and radiographically consistent with mild encephalitis/encephalopathy with reversible splenial lesion (MERS). He improved with supportive therapy with intravenous glucose and vitamins, and respiratory support was not required after 2 days. Neurological signs of ptosis, strabismus and ataxia persisted for a few months before recovering, and the MRI lesions resolved after 4 months (not shown).

At age 5 6/12 years, following diarrhea, fever and fasting for 24 hours, he presented again encephalopathic with decreased consciousness (GCS 10), mild hypoglycemia 56 mg/dL, metabolic acidosis (pH 7.3, bicarbonate 14.2 mmol/L), elevated serum lactate 6.5 mmol/L, and in urine elevated lactate, ketones, and dicarboxylic acids. Plasma ammonia and muscle enzymes remained normal. He recovered again with intravenous glucose administration. He was stable for several years but had learning difficulties and unilateral hearing loss.

At age 13 6/12 years, he had headache, asthenia and anorexia for 36 hours, and became stuporous without metabolic abnormalities. Brain MRI demonstrated hyperintense lesions on T2-weighted imaging of the caudate nuclei and putamina (Figure 3H-I, white arrowheads), and the ventromedial aspects of the thalami (Figure 3I, white notched arrowheads). Clinically, he recovered completely within 2 weeks. He has remained stable and generally healthy, except for easy fatigue and learning difficulties with memory and attention deficits.

A fasting test performed outside an acute episode showed normal ketogenesis. Fibroblasts showed normal enzyme activities of pyruvate carboxylase and succinyl-CoA:3-keto acid coenzyme A transferase (SCOT), and leucocytes had normal pyruvate dehydrogenase activity. Molecular analysis of *SLC19A3* and *DLD* was normal. Methemoglobin levels were borderline elevated 2.1% and 2.9% (normal <2%).

### 3.2 | Mitochondrial functional studies

In subject A.II-3, analysis of the respiratory chain enzyme complexes during an acute episode showed markedly reduced complex IV activity with mildly decreased complex II activity, more so in muscle than in liver (Table 1). Surprisingly, BN PAGE analysis in muscle and liver showed only a mild decrease in complex IV activity within the range of normal variation (Figure 4A), suggesting that it affected enzymatic activity more than the structure or amount of the complex. In fibroblasts, analysis of mitochondrial function was not possible as cells from subject A.II-3 rapidly died on initial culture of



**TABLE 1** Respiratory chain enzyme activities in patient A.II.3

	Activity (controls) nmol min <sup>-1</sup> mg protein <sup>-1</sup>	% of normal	SD	Activity/CS	SD	Activity/CO II	SD
Muscle							
Complex I	44.2 (23.6-74.8)	98%	0.1	176 (98-271)	0.1	888 (285-767)	1.9
Complex II	49.8 (49.0-133.4)	56%	-1.8	198 (251-573)	-1.8	NA	NA
Complex III	20.3 (5.7-31.4)	117%	0.5	81 (19-72)	0.5	408 (45-369)	1.2
Complex II-III	45.7 (34.2-107.6)	72%	-0.7	182 (172-472)	-0.1	918 (549-1226)	0.7
Complex IV	<b>0.4</b> (1.1-3.8)	<b>15%</b>	<b>-4.3</b>	<b>2</b> (4-23)	<b>-4.3</b>	<b>8</b> (11-68)	<b>-4.6</b>
CS	251.7 (159.8-353.3)	98%	0.0	NA	NA	NA	NA
Liver							
Complex I	60.8 (14.4-56.0)	172%	1.5	600 (162-730)	0.9	264 (68-252)	1.3
Complex II	230.1 (174.7-309.8)	99%	0.1	2269 (2304-3311)	-1.3	NA	NA
Complex III	27.6 (13.8-27.6)	145%	3.1	273 (128-315)	0.6	120 (50-118)	1.0
Complex II-III	50.0 (10.8-107.3)	85%	0.4	493 (138-1062)	-0.2	217 (62-383)	0.0
Complex IV	<b>0.4</b> (0.5-3.2)	<b>22%</b>	-1.8	<b>4</b> (6-35)	<b>-2.4</b>	<b>2</b> (3-13)	<b>-2.5</b>
CS	101.4 (59.5-109.3)	120%	0.1	NA	NA	NA	NA

Note: The activities in muscle and liver tissue of each respiratory chain complex and of the combined complex II-III are shown expressed as nmol min<sup>-1</sup> mg protein<sup>-1</sup> and as a ratio of the activity over citrate synthase activity (activity/CS) and as a ratio over the activity of complex II (activity/Co II). The patient's values are followed by normal values in parentheses and the percentage of the average normal value is provided. The values are also expressed as standard deviations (Z-score) of the log transformed values of controls, which are normally distributed. Activities that are reduced are highlighted in bold.

Abbreviations: CS, citrate synthase; CO II, complex II.

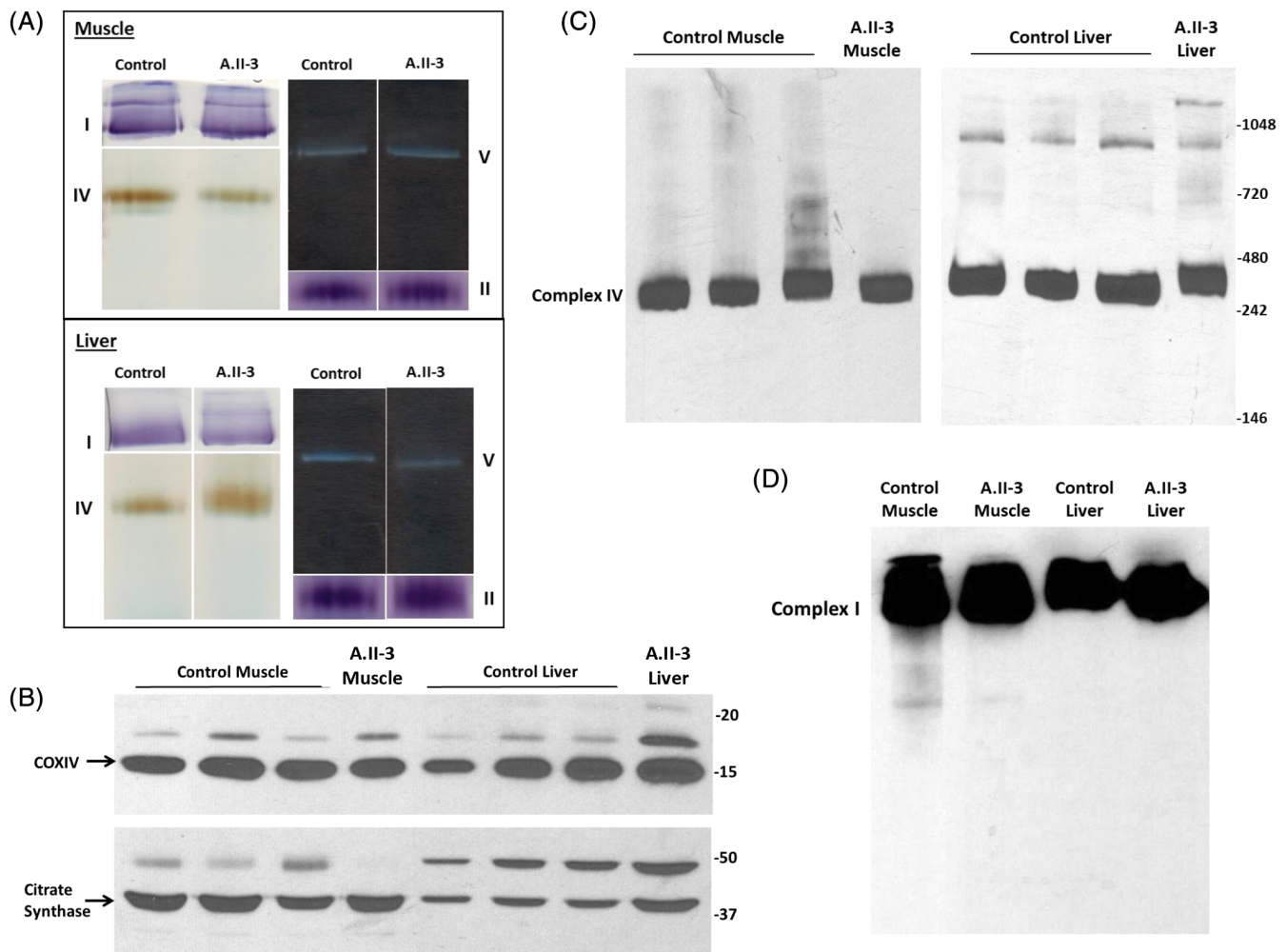
unknown cause. Fibroblasts from subject B.II-3 grew well and both respiratory chain enzyme activities and BN PAGE with in-gel activity staining were normal (Supplemental Table S3). We next evaluated directly if the reduced activity evident in muscle and liver of subject A.II-3 was due to a decrease in complex IV assembly. Despite the strong decrease in complex IV activity, the amount of a protein COXIV, as a representative of complex IV subunits, was normal (Figure 4B). Following separation on a nondenaturing gel, normal amounts of fully assembled complex IV were present in both muscle and liver in the patient compared to three controls (Figure 4C). The integrity of complex I assessed by the amount of NDUFB8 as a marker subunit protein was normal, as was its assembly assessed on a nondenaturing gel (Figure 4D). Thus, the marked decrease in complex IV activity was not caused by a problem in complex IV biogenesis but rather its function.

### 3.3 | Genetic studies

The *SQOR* gene has 8 transcripts. The main protein coding transcript *SQOR*-201 has 10 exons. The transcript has 1701 bp and encodes a protein of 450 residues (Uniprot Q9Y6N5). Whole exome studies in the proband A.II-3 showed a homozygous missense variant NM\_021199.4:

c.637G > A, p.(Glu213Lys) in exon 5 of the *SQOR* gene, genomic coordinates: hg19:chr15:45965982G > A, which was confirmed by Sanger sequencing (Figure 2C). Her sister A.II-2 was also homozygous for this variant and both parents and two unaffected siblings were heterozygous carriers (Figure 2A). Mutation Taster predicted this variant to be disease-causing, Polyphen2 rated it as "possibly damaging" (score 0.915), and the scaled CADD score was 34 (raw score 5.03). The *SQOR* p.Glu213Lys variant is not reported in 1000 genomes or ExAC. In gnomAD it was present in only one non-Finnish European allele (minor allele frequency [MAF] in this population  $0.9 \times 10^{-5}$ , MAF  $0.4 \times 10^{-5}$  in all populations). Phylogenetically, in most species, this amino acid is an acidic residue, either a glutamic acid or an aspartic acid; the mutation changes Glu213 to a basic lysine.

Whole exome sequencing in subject B.II-3 revealed a homozygous 1 bp deletion c.446delT, p.(Leu149Argfs\*18), genomic coordinates chr15:45962165delT, in exon 3 of the *SQOR* gene, which was confirmed by Sanger sequencing (Figure 2C). This change predicts a loss of *SQOR* protein function possibly by nonsense-mediated decay of the mutant RNA or by introducing a frameshift leading to a premature truncation of protein translation. This variant was present in heterozygous state in gnomAD only once (MAF in this population  $3.98 \times 10^{-6}$ ). Sanger sequencing



**FIGURE 4** Assembly and activity of mitochondrial complexes. A, Mitochondrial complexes were separated on a blue native (BN) gel and assayed with in-gel activity staining in muscle (upper panel) and in liver (lower panel). A mild decrease in complex IV staining was observed in muscle for patient A.II-3. B, In muscle and liver of patient A.II-3 a normal amount of the complex IV subunit COXIV is seen, using citrate synthase as a loading control. C, Both muscle and liver exhibit normal levels of fully assembled complex IV in patient A.II-3 compared to control samples as revealed by blue gel followed by western blotting and detection with COXIV antibody. D, The assembly of complex I is normal in patient A.II-3 compared to controls as identified by BN gel followed by western blotting and detection with an antibody against NDUFS2

revealed this change also in a homozygous state in the asymptomatic sisters B.II-1 and B.II-2 (Figure 2B).

### 3.4 | SQOR functional studies

To understand the impact of the mutation on the protein, the missense variant was mapped on the crystal structure of human SQOR (Figure 2D). In the structure, Glu-213 is remote from the active site, but resides in an  $\alpha$ -helix and is proximal to Arg-217 on the same helix and Arg-222 on an adjacent  $\alpha$ -helix. The side chain of Glu213 is predicted to be involved in hydrogen bonding interactions with the side chains of these two arginine residues (Figure 2D). Introduction of the positively charged lysine in the

Glu213Lys mutant is predicted to disrupt these electrostatic interactions and destabilize the tertiary structure of the protein. This structure suggests that the Glu213Lys variant is likely to impair protein stability rather than affect the catalytic mechanism.

Indeed, attempts to express the recombinant Glu213Lys human SQOR protein in *E. coli* using the methods employed for wild type proteins<sup>17</sup> as well as several modifications to growth and purification conditions, repeatedly resulted in the majority of the recombinant protein aggregating in inclusion bodies. The yield of soluble Glu213Lys SQOR protein was very low and the protein appeared to be devoid of FAD as assessed by the lack of an associated ultraviolet-visible absorption spectrum.

Western blot analysis showed very low levels of SQOR protein in muscle and liver tissues of subject A.II-3 and in fibroblasts of subject B.II-3 each compared to three controls (Figure 2E).

To evaluate the functional impact, an SQOR enzyme activity assay using sulfite as acceptor was developed using liver tissue mitochondrial membrane fractions. The initial rate was linearly dependent on lysate protein concentration over the range of 0.064 to 0.698 mg protein/mL (final concentration) ( $r^2 = .98$ ) (Supplemental Figure S1A). A  $K_M$  of  $2.2 \pm 1.1$  mM for sulfide and a  $V_{max}$  of  $27.8 \pm 8.9$  nmol min<sup>-1</sup> mg protein<sup>-1</sup> at 37°C were observed (Supplemental Figure S1C). The  $V_{max}$  value was obtained at sulfite concentrations of around 1 to 2 mM (Supplemental Figure S1D), and at coenzyme Q<sub>1</sub> concentrations of 100 μM (Supplemental Figure S1E). The coefficient of variation (COV) in the same liver tissue sample was 14%. SQOR activity in subject A.II-3 liver was severely reduced to 6% of the average of control values ( $0.71 \pm 0.57$  vs  $11.42 \pm 3.75$  nmol min<sup>-1</sup> mg protein<sup>-1</sup>, normal range 7.56-16.94;  $n = 5$ ) (Supplementary Figure S2A).

To facilitate future diagnosis of SQOR deficiency, the enzyme assay was modified for use with fibroblasts and PBMCs. In fibroblasts, the initial rate of enzyme activity was linear over a lysate protein concentration range from 0.01 to 0.34 mg protein mL<sup>-1</sup> ( $r^2 = .98$ ) (Supplemental Figure S1B). With control fibroblasts, a specific activity of  $82.1 \pm 21.2$  nmol min<sup>-1</sup> mg protein<sup>-1</sup> (normal range 51.6-115.2;  $n = 12$ ), COV 26% was obtained, whereas subject B.II-3 fibroblasts had SQOR activity of  $0.41 \pm 0.16$  nmol min<sup>-1</sup> mg protein<sup>-1</sup>, 0.49% of average control value (Supplemental Figure S2B). Analysis of control PBMCs yielded a specific activity of  $43.6 \pm 2.1$  nmol min<sup>-1</sup> mg protein<sup>-1</sup> ( $n = 8$ ), COV 5%. Additional family members who were found to be carriers for this mutation had 62%, 66%, and 42% of mean control enzyme activity in PBMCs.

The homozygosity of the patient and the identification of additional carriers in distant family members suggested that Glu213Lys may represent a founder mutation. Targeted mutation screening of de-identified DNA samples of 40 subjects from this population identified three carriers (carrier frequency  $P = .08 \pm .04$ ), implying an affected incidence of 1 in 711. Untargeted population analysis of the gnomAD database using predictive software estimated a carrier frequency of  $P = .0023$  (1 in 215), implying an incidence of 1 in 184 477 for biallelic pathogenic variants.

## 4 | DISCUSSION

The Glu213Lys variant replaces an acidic with a basic amino acid disrupting hydrogen bonding interactions

and destabilizing the SQOR protein. This protein instability resulted in strongly reduced protein levels and severely decreased enzyme activity, but not its complete absence. The protein instability severely limited bacterial expression of the recombinant Glu213Lys human SQOR enzyme. These functional data confirm bioinformatic predictions that this variant is pathogenic. It tracks with deficient enzyme activity in the family and represents a founder variant in the Lehrerleut Hutterite population with a carrier frequency estimated to be as high as 1 in 13 based on limited screening of samples from this population. The frameshift mutation in Family B likewise resulted in severely reduced SQOR protein and enzyme activity. These pathogenic variants limited the ability of the patient to metabolize H<sub>2</sub>S, and presumably resulted in episodic accumulation.

In all patients, acute symptoms were triggered by infections and fasting, episodes associated with protein catabolism. Outside acute events, our patients generally appeared healthy, and likewise two siblings in Family B have remained asymptomatic for prolonged time. They did not exhibit symptoms such as chronic gastrointestinal upset or a tendency for petechiae that have previously been attributed to excess H<sub>2</sub>S in patients with pathogenic variants in *ETHE1* resulting in deficient persulfide dioxygenase activity, the next step in the pathway.<sup>4</sup> In contrast to *ETHE1* deficiency, ethylmalonic acid was not noted in urine organic acid analysis, supporting that the inhibition of short chain acyl-CoA dehydrogenase could be due to accumulating coenzyme A-persulfide in the intact SQOR protein in *ETFE1* deficiency, but not in SQOR deficiency.<sup>27</sup> Chronic loss of complex IV protein attributed to excess H<sub>2</sub>S in the mouse model of *ETHE1* was also not observed in our patient.<sup>5</sup> Rather, a functional deficiency of complex IV activity was seen, while the levels and assembly of complex IV appeared comparable to controls. Deficient complex IV activity results in an acute energy defect which in brain usually presents as Leigh disease,<sup>21</sup> although some genetic causes also include myopathy or cardiomyopathy (eg, *SCO2*).<sup>30</sup> In our patients, the cardiovascular system appeared to be relatively spared with cardiovascular symptoms limited to transient hypotension and shock present only in the context of multiorgan failure, which recovered with supportive care. Acute H<sub>2</sub>S toxicity causes depression in cardiac contractility and pulseless electrical activity.<sup>31</sup> In an animal model of acute H<sub>2</sub>S intoxication leading to 75% mortality, 25% of surviving animals had neurological lesions of neuronal necrosis of the cortex, putamen and thalamus, which closely resembles the symptoms noted in patient A.II-2.<sup>31</sup> The striking involvement of the hippocampus and mammillary body in patient A.II-3 without prolonged seizures likely represents the SQOR deficiency

rather than an injury secondary to her poor state. In mouse, SQOR protein is present in the hippocampus (Maclean, unpublished observation<sup>25</sup>). Given the long asymptomatic period in subject A.II-2 and subjects B.II-1 and B.II-2, we consider it possible that undiagnosed patients exist, including in the Hutterite population.

The presence of normal amounts of cytochrome c oxidase complex but severely reduced activity indicates that the fundamental pathophysiological mechanism of SQOR deficiency appears to be due to the resulting accumulation of toxic metabolites, rather than disruption of complex IV biogenesis. Acute illness is associated with catabolism and associated endogenous protein breakdown. The resulting increase in cysteine can increase H<sub>2</sub>S generation, which would accumulate in the absence of sufficient SQOR enzyme activity. At high levels, H<sub>2</sub>S can coordinate to the iron in the heme a group of complex IV resulting in strong inhibition.<sup>3</sup> The complete absence of SQOR in yeast impaired growth under obligate aerobic conditions, whereas mutants that retain low levels of activity restored oxidative growth, consistent with the detoxifying role attributed to SQOR.<sup>32</sup> We hypothesize that in our patients, the low residual SQOR activity was likely sufficient to support H<sub>2</sub>S clearance during stable episodes but was insufficient during acute illness with catabolism resulting in H<sub>2</sub>S accumulation and toxicity.

During acute H<sub>2</sub>S intoxication, two pools accumulate: a soluble and diffusible pool of H<sub>2</sub>S gas in equilibrium with the sulfide (HS<sup>-</sup>) anion, and a pool of sulfide bound to metalloproteins (eg, hemes) and cysteine residues in a variety of proteins.<sup>31, 33</sup> It is unclear to what extent these cysteine adducts can form nonenzymatically in the absence of functional SQOR. The toxicity mechanism in our patients might be partially different from that of acute exogenous H<sub>2</sub>S intoxication. In our patient A.II-3, the more severe reduction in complex IV activity was noted in muscle tissue, where the sulfide generating enzymes could not be detected, compared to liver tissue where the enzymes were present and unchanged vs controls (Supplemental Figure S2). This illustrates the importance of great diffusibility of H<sub>2</sub>S, which is more soluble and diffusible than oxygen or carbon dioxide.<sup>31</sup> A large pool of H<sub>2</sub>S bound to ferrous myoglobin has been proposed to explain the sensitivity of muscle tissue to H<sub>2</sub>S toxicity.<sup>34</sup> SQOR has a wide tissue distribution including in neurons, oligodendrocytes and vascular endothelial cells in brain, likely reflecting a need to protect against this toxin that readily diffuses across membranes.<sup>35, 36</sup> Rapid diffusion of H<sub>2</sub>S from tissue culture medium can explain the normal complex IV activity in cultured fibroblasts.

Recognition of the enzymatic basis and intoxication of the respiratory chain in the pathophysiology of SQOR deficiency opens up possible therapeutic opportunities. First,

reduction in H<sub>2</sub>S production should be pursued. Since the majority of H<sub>2</sub>S is generated from cysteine catabolism in the body and gut microbiome,<sup>13</sup> limiting dietary cysteine and aggressive prevention of episodes triggering catabolism by adequate provision of calories and avoidance of prolonged fasting can be considered. Catabolic episodes likely triggered the acute crisis in our patients. H<sub>2</sub>S is produced by  $\gamma$ -proteobacteria including certain gram-negative and *Clostridia* species, which can be increased by omeprazole or lansoprazole,<sup>14</sup> and can be therapeutically reduced by treatment with metronidazole, similar to the treatment of ETHE1 disorder.<sup>37</sup> Certain vegetables, such as garlic, onions, and certain cruciferous vegetables contain organic polysulfides, which can directly release H<sub>2</sub>S and could be avoided.<sup>14</sup> Next, the toxic effects of accumulated H<sub>2</sub>S could be mitigated. Toxicological studies have shown that hydroxocobalamin can bind H<sub>2</sub>S and pharmacological doses have been used for detoxification during acute H<sub>2</sub>S poisoning, similar to the treatment of cyanide which shares a similar pathophysiology.<sup>31, 33, 38</sup> During acute symptomatic periods in illness, acute administration of very high doses of hydroxocobalamin (70 mg/kg) may be required. Given the rapidity of onset of life-threatening events, which in our patients occurred within hours, preventive daily subcutaneous administration of pharmacologic doses of hydroxocobalamin, as is used to treat cobalamin C disorder, should be considered. Further, methylene blue has been shown to reduce mortality due to acute H<sub>2</sub>S poisoning by reducing its mitochondrial toxicity.<sup>31, 39</sup>

In summary, we describe a new genetic cause of Leigh disease associated with pathogenic variants in SQOR, whereby intermittent accumulation of H<sub>2</sub>S results in acute isolated complex IV deficiency. We developed an enzyme assay that allows noninvasive measurement of the enzyme activity, which is useful for diagnosis. Recognition of SQOR deficiency as a cause of Leigh diseases is clinically important since it suggests several treatment modalities to prevent neurological complications.

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#### CONFLICT OF INTERESTS

J. V. H. and A. A. L. participate in clinical trials of mitochondrial disorders by Stealth Biotherapeutics, Inc. All other authors deny any real or apparent conflict of interest in the field of mitochondrial diseases.

#### AUTHOR CONTRIBUTIONS

Study concept and design: Johan L. K. Van Hove and Marisa W. Friederich. Patient care data: Johan L. K. Van Hove, Abdallah F. Elias, Alice Kuster, Austin A. Larson, Jaclyn Haven, David M. Mirsky, Louise Goujon, Thomas Lefrancois. Molecular data: Marisa W. Friederich, Abdallah F. Elias, Lucia Laugwitz, David Dimmock, Katie Styren, Jonathan Schoof, Maike Friederich, Curtis R. Coughlin, Tobias B. Haack, Johan L. K. Van Hove. Laboratory studies: Marisa W. Friederich, Aaron P. Landry, Logan Ellwood-Digel, Hua Jiang, Kenneth N. MacLean, Ruma Banerjee. First draft writing: Marisa W. Friederich, Johan L. K. Van Hove. Critical rewriting: Marisa W. Friederich, Abdallah F. Elias, David M. Mirsky, David Dimmock, Kenneth N. MacLean, Curtis R. Coughlin, Ruma Banerjee, Tobias B. Haack, Johan L. K. Van Hove. Leadership and funding: Marisa W. Friederich, Abdallah F. Elias, Kenneth N. MacLean, Ruma Banerjee, Tobias B. Haack, Johan L. K. Van Hove. Final responsibility and guarantor: Johan L. K. Van Hove.

#### REFERENCES

- Kolluru GK, Shen X, Bir SC, Kevil CG. Hydrogen sulfide chemical biology: pathophysiological roles and detection. *Nitric Oxide*. 2013;35:5-20.
- Kabil O, Vitvitsky V, Banerjee R. Sulfur as a signaling nutrient through hydrogen sulfide. *Annu Rev Nutr*. 2014;34:171-205.
- Cooper CE, Brown GC. The inhibition of mitochondrial cytochrome oxidase by the gases carbon monoxide, nitric oxide, hydrogen cyanide and hydrogen sulfide: chemical mechanism and physiological significance. *J Bioenerg Biomembr*. 2008;40:533-539.
- Tiranti V, Viscomi C, Hildebrandt T, et al. Loss of ETHE1, a mitochondrial dioxygenase, causes fatal sulfide toxicity in ethylmalonic encephalopathy. *Nat Med*. 2009;15:200-205.
- Di Meo I, Fagiolaro G, Prella A, Viscomi C, Zeviani M, Tiranti V. Chronic exposure to sulfide causes accelerated degradation of cytochrome c oxidase in ethylmalonic encephalopathy. *Antiox Redox Sign*. 2011;15:353-362.
- Hildebrandt TM, Di Meo I, Zeviani M, Viscomi C, Braun H-P. Proteome adaptations in Ethe1-deficient mice indicate a role in lipid catabolism and cytoskeleton organization via post-translational modification. *Biosci Rep*. 2013;33:575-584.
- Di Meo I, Lamperti C, Tiranti V. Mitochondrial diseases caused by toxic compound accumulation: from etiopathology to therapeutic approaches. *EMBO Mol Med*. 2015;7:1257-1266.
- Luna-Sánchez M, Hidalgo-Gutiérrez A, Hildebrandt TM, et al. CoQ deficiency causes disruption of mitochondrial sulfide oxidation, a new pathomechanism associated with this syndrome. *EMBO Mol Med*. 2017;9(1):78-95.
- Ziosi M, Di Meo I, Kleiner G, et al. Coenzyme Q deficiency causes impairment of the sulfide oxidation pathway. *EMBO J*. 2017;9(1):96-111.
- Chiku T, Padovani D, Zhu W, Singh S, Vitvitsky V, Banerjee R. H<sub>2</sub>S biogenesis by human cystathionine  $\gamma$ -lyase leads to the novel sulfur metabolites lanthionine and homolanthionine and is responsive to the grade of hyperhomocysteinemia. *J Biol Chem*. 2009;284:11601-11612.
- Singh S, Padovani D, Leslie RA, Chiku T, Banerjee R. Relative contributions of cystathionine  $\beta$ -synthase and  $\gamma$ -cystathioninase to H<sub>2</sub>S biogenesis via alternative trans-sulfuration reactions. *J Biol Chem*. 2009;284:22457-22466.
- Yin J, Ren W, Yang G, et al. L-cysteine metabolism and its nutritional implications. *Mol Nutr Food Res*. 2016;60:134-146.
- Magee EA, Richardson CJ, Hughes R, Cummings JH. Contribution of dietary protein to sulfide production in the large intestine: an in vitro and a controlled feeding study in humans. *Am J Clin Nutr*. 2000;72:1488-1494.
- Wallace JL, Motta JP, Buret AG. Hydrogen sulfide: an agent of stability at the microbiome-mucosa interface. *Am J Physiol Gastrointest Liver Physiol*. 2018;314:G143-G149.
- Libiad M, Vitvitsky V, Bostelaar T, et al. Hydrogen sulfide perturbs mitochondrial bioenergetics and triggers metabolic reprogramming in colon cells. *J Biol Chem*. 2019;294(13):12077-12090.
- Jackson MR, Melideo SL, Jorns MS. Human sulfide:quinone oxidoreductase catalyzes the first step in hydrogen sulfide metabolism and produces a sulfane sulfur metabolite. *Biochemistry*. 2012;51:6804-6815.
- Libiad M, Yadav PK, Vitvitsky V, Martinov M, Banerjee R. Organization of the human mitochondrial hydrogen sulfide oxidation pathway. *J Biol Chem*. 2014;289:30901-30910.
- Landry AP, Ballou DP, Banerjee R. H<sub>2</sub>S oxidation by nanodisc-embedded human sulfide quinone oxidoreductase. *J Biol Chem*. 2017;292:11641-11649.

19. Rahman S, Blok RB, Dahl HH, et al. Leigh syndrome: clinical features and biochemical and DNA abnormalities. *Ann Neurol*. 1996;39(3):343-351.
20. Lake NJ, Compton AG, Rahman S, Thorburn DR. Leigh syndrome: one disorder, more than 75 monogenic causes. *Ann Neurol*. 2016;79(2):190-203.
21. DiMauro S, Servidei S, Zeviani M, et al. Cytochrome c oxidase deficiency in Leigh syndrome. *Ann Neurol*. 1987;22:498-506.
22. Karczewski KJ, Francioli LC, Tiao G, et al. Variation across 141,456 human exomes and genomes reveals the spectrum of loss-of-function intolerance across human protein-coding genes. *BioRxiv*. 2019. <https://doi.org/10.1101/531210>
23. Chatfield KC, Coughlin CR 2nd, Friederich MW, et al. Mitochondrial energy failure in HSD10 disease is due to defective mtDNA transcript processing. *Mitochondrion*. 2015;21:1-10.
24. Friederich MW, Erdogan AJ, Coughlin CR II, et al. Mutations in the accessory subunit NDUFB10 result in isolated complex I deficiency and illustrate the critical role of intermembrane space import for complex I holoenzyme assembly. *Hum Mol Genet*. 2017;26(4):702-716.
25. Maclean KN, Jiang H, Greiner LS, Allen RH, Stabler SP. Long-term betaine therapy in a murine model of cystathionine beta-synthase deficient homocystinuria: decreased efficacy over time reveals a significant threshold effect between elevated homocysteine and thrombotic risk. *Mol Genet Metab*. 2012;105(3):395-403.
26. Banerjee R, Chiku T, Kabil O, Libiad M, Motl N, Parmod K. Assay methods for H<sub>2</sub>S biogenesis and catabolism enzymes. *Methods Enzymol*. 2015;554:189-200.
27. Landry AP, Moon S, Kim H, et al. A catalytic trisulfide quinone oxidoreductase catalyzes coenzyme A persulfide synthesis and inhibits butyrate oxidation. *Cell Chem Biol*. 2019;26:1515-1525.
28. Schrödinger, LLC. The PyMOL Molecular Graphics System, Version 2.0, <https://pymol.org>; 2019.
29. Zhang W, Cui H, Wong LJ. Comprehensive one-step molecular analyses of mitochondrial genome by massively parallel sequencing. *Clin Chem*. 2012;58(9):1322-1331.
30. Rak M, Bénit P, Chrétien D, et al. Mitochondrial c oxidase deficiency. *Clin Sci*. 2016;130(6):393-407.
31. Haouzi P, Sonobe T, Judenherc-Haouzi A. Developing effective countermeasures against acute hydrogen sulfide intoxication: challenges and limitations. *Ann N Y Acad Sci*. 2016;1374:29-40.
32. Vande Weghe JG, Ow DW. A fission yeast gene for mitochondrial sulfide oxidation. *J Biol Chem*. 1999;274:13250-13257.
33. Haouzi P, Sonobe T, Torsell-Tubbs N, Prokopczyk B, Chenuel B, Klingerman CM. *In vivo* interactions between cobalt and ferric compounds and the pools of sulphide in the blood during and after H<sub>2</sub>S poisoning. *Toxicol Sci*. 2014;141(2):493-504.
34. Bostelaar T, Vitvitsky V, Kumutima J, et al. Hydrogen sulfide oxidation by myoglobin. *J Am Chem Soc*. 2016;138(27):8476-8488.
35. Ackermann M, Kubitzka M, Hauska G, Piña AL. The vertebrate homologue of sulfide-quinone reductase in mammalian mitochondria. *Cell Tissue Res*. 2014;358:779-792.
36. Ackermann M, Kubitzka M, Brawanski A, Hauska G, Piña AL. The vertebrate homolog of sulfide-quinone reductase is expressed in mitochondria of neuronal tissues. *Neuroscience*. 2011;199:1-12.
37. Viscomi C, Burlina AB, Dweikat I, et al. Combined treatment with oral metronidazole and N-acetylcysteine is effective in ethylmalonic encephalopathy. *Nat Med*. 2010;16(8):869-871.
38. Haouzi P, Chenuel B, Sonobe T. High-dose hydroxocobalamin administered after H<sub>2</sub>S exposure counteracts sulfide-poisoning-induced cardiac depression in sheep. *Clin Toxicol*. 2015;53:28-36.
39. Houzi P, Tubbs N, Chaeung J, Judenher-Haouzi A. Methylene blue administration during and after life-threatening intoxication by hydrogen sulfide: efficacy studies in adult sheep and mechanisms of action. *Toxicol Sci*. 2019;168(2):443-459.

## SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of this article.

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