Secondary mitochondrial dysfunction in propionic aciduria: a pathogenic role for endogenous mitochondrial toxins

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Mitochondrial dysfunction during acute metabolic crises is considered an important pathomechanism in inherited disorders of propionate metabolism, i.e. propionic and methylmalonic acidurias. Biochemically, these disorders are characterized by accumulation of propionyl-CoA and metabolites of alternative propionate oxidation. In the present study, we demonstrate uncompetitive inhibition of PDHc (pyruvate dehydrogenase complex) by propionyl-CoA in purified porcine enzyme and in submitochondrial particles from bovine heart being in the same range as the inhibition induced by acetyl-CoA, the physiological product and known inhibitor of PDHc. Evaluation of similar range as the inhibition induced by acetyl-CoA, the physiological metabolism and ammonia detoxification [1]. Clinically, PA is characterized by an accumulation of propionate, 3-hydroxypropionyl-CoA, subsequently being fed as anaplerotic precursor into the tricarboxylic acid cycle. Inherited deficiency of propionyl-CoA carboxylase, i.e. PA (propionic aciduria), is biochemically characterized by an accumulation of propionate, 3-hydroxypropionate, 2-methylcitrate and propionylglycine, demonstrating alternative propionate oxidation in these patients. Furthermore, metabolic acidosis, ketosis, increased lactate concentrations, hypoglycaemia and hyperammonaemia are found during metabolic derangement, highlighting secondarily compromised energy metabolism and ammonia detoxification [1]. Clinically, PA is complicated by acute life-threatening metabolic crises, which are precipitated by catabolic state and result in multiple organ failure or even death if untreated [1]. Like other organic acidurias (e.g. methylmalonic aciduria), age at clinical onset and disease course may vary between individual patients presenting with neonatal metabolic encephalopathy (neonatal onset), recurrent episodes of ketoacidotic coma or Reye-like syndromes (chronic intermittent form), or psychomotor retardation and failure to thrive in the absence of acute crises (chronic progressive form). Similarly to neonatal onset of PA patients, PCCA−/− mice, a transgenic mouse model for PA, develop lethal ketoacidosis shortly after birth [2].

Despite improvements in the diagnostic work-up and management of affected patients during the last decades, the neurological outcome remains disappointing in PA patients [1,3]. In particular, mental retardation and movement disorders secondary to cortical atrophy, leucoencephalopathy and basal ganglia injury are still frequently found even in early diagnosed children. The biochemical abnormalities found during acute metabolic crises have led to the suggestion that impairment of OXPHOS (oxidative phosphorylation) is crucial for the pathogenesis of this disease. Notably, accumulating metabolites of alternative propionate oxidation have been suggested to act as endogenous inhibitors of energy metabolism in different in vitro models [4–7]; however, the pathophysiological impact of these findings on PA still remains unclear. Here, we report severe disturbance of mitochondrial energy metabolism in muscle tissues from two PA patients and demonstrate in vitro that propionyl-CoA-induced mitochondrial dysfunction plays a central role in this scenario.

INTRODUCTION

Propionyl-CoA is an intermediate metabolite in the final common catabolic pathways of the amino acids L-isoleucine, L-methionine, L-threonine and L-valine as well as odd-chain fatty acids and cholesterol. It is carboxylated by propionyl-CoA carboxylase (EC 6.4.1.3) to methylmalonyl-CoA and then is converted by methylmalonyl-CoA mutase (EC 5.4.99.2) into succinyl-CoA, subsequently being fed as anaplerotic precursor into the tricarboxylic acid cycle. Inherited deficiency of propionyl-CoA carboxylase, i.e. PA (propionic aciduria), is biochemically characterized by an accumulation of propionate, 3-hydroxypropionate, 2-methylcitrate and propionylglycine, demonstrating alternative propionate oxidation in these patients. Furthermore, metabolic acidosis, ketosis, increased lactate concentrations, hypoglycaemia and hyperammonaemia are found during metabolic derangement, highlighting secondarily compromised energy metabolism and ammonia detoxification [1]. Clinically, PA is complicated by acute life-threatening metabolic crises, which are precipitated by catabolic state and result in multiple organ failure or even death if untreated [1]. Like other organic acidurias (e.g. methylmalonic aciduria), age at clinical onset and disease course may vary between individual patients presenting with neonatal metabolic encephalopathy (neonatal onset), recurrent episodes of ketoacidotic coma or Reye-like syndromes (chronic intermittent form), or psychomotor retardation and failure to thrive in the absence of acute crises (chronic progressive form). Similarly to neonatal onset of PA patients, PCCA−/− mice, a transgenic mouse model for PA, develop lethal ketoacidosis shortly after birth [2]. Despite improvements in the diagnostic work-up and management of affected patients during the last decades, the neurological outcome remains disappointing in PA patients [1,3]. In particular, mental retardation and movement disorders secondary to cortical atrophy, leucoencephalopathy and basal ganglia injury are still frequently found even in early diagnosed children. The biochemical abnormalities found during acute metabolic crises have led to the suggestion that impairment of OXPHOS (oxidative phosphorylation) is crucial for the pathogenesis of this disease. Notably, accumulating metabolites of alternative propionate oxidation have been suggested to act as endogenous inhibitors of energy metabolism in different in vitro models [4–7]; however, the pathophysiological impact of these findings on PA still remains unclear. Here, we report severe disturbance of mitochondrial energy metabolism in muscle tissues from two PA patients and demonstrate in vitro that propionyl-CoA-induced mitochondrial dysfunction plays a central role in this scenario.

Abbreviations used: BN-PAGE, blue native PAGE; E1, pyruvate decarboxylase; E3, dihydrolipoyl dehydrogenase; KGDHc, α-ketoglutarate dehydrogenase complex; mDNA, mitochondrial DNA; ND6, NADH dehydrogenase 6; OXPHOS, oxidative phosphorylation; PA, propionic aciduria; PDHc, pyruvate dehydrogenase complex; SMP, submitochondrial particle.

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**EXPERIMENTAL**

**Patient 1**

This girl was born at term as the second child of non-consanguineous Caucasian parents. At the third day of life, she was admitted because of progressive feeding refusal, lethargy and abnormal breathing. Laboratory investigations revealed a severe metabolic acidosis [pH 7.01; pCO\(_2\) 18 mmHg (1 mmHg = 0.133 kPa); bicarbonate 5 mM; base excess 25 mM] and hyperammonaemia (830 \(\mu\)M; normal < 80 \(\mu\)M). The amino acid analyses in plasma revealed no abnormalities. Analysis of urine organic acids by GC/MS showed biochemical abnormalities characteristic for PA with increased concentrations of \(\beta\)-hydroxybutyric acid (6373 mmol/mol of creatinine; control range: 0–45 mmol/mol of creatinine), \(\beta\)-hydroxypropionic acid (21461 mmol/mol of creatinine; control range: 0–160 mmol/mol of creatinine), \(\beta\)-hydroxyisovaleric acid (176 mmol/mol of creatinine; control range: 0–10 mmol/mol of creatinine), lactic acid (776 mmol/mol of creatinine; control range: 0–270 mmol/mol of creatinine), propionylglycine, methylcitrate and tiglylglycine. The last three metabolites varying between 40 and 80 \(\mu\)M; control range: 0–270 mmol/mol of creatinine, propionyl-

Glycine, methylcitrate and tiglylglycine. The last three metabolites varying between 40 and 80 \(\mu\)M; control range: 0–270 mmol/mol of creatinine, propionyl-

Extracorporal detoxification was performed. Besides a high caloric intake and transient stop of protein intake, metronidazol, 

Repeatedly showed increased concentrations of propionylcarnitine and the only abnormal clinical sign at that time was a slight axial hypotonia. Maintenance treatment consisted of a natural protein intake (1.0 g/kg per day) supplemented with an amino acid mixture lacking precursor amino acids of propionyl-CoA (i.e. L-isoleucine, L-methionine, L-threonine and L-valine) and application of L-carnitine and biotin. Application of propionyl-CoA carboxylase activity in leucocytes showed deficient activity of this enzyme (0.1 nmol h\(^{-1}\) · mg of protein\(^{-1}\); control: 2.3 nmol h\(^{-1}\) · mg of protein\(^{-1}\)) and thus confirmed the diagnosis of PA. She was admitted to hospital numerous times due to metabolic derangements, the last one, at age 5 years, being fatal. The child had a severe myopathic appearance during life, which was, in combination with repeatedly increased blood lactate concentrations (2.0–7.2 mM; normal < 2.1 mM), a reason to perform a muscle biopsy for mitochondrial studies.

**Patient 2**

This boy was born at term following an uncomplicated pregnancy and delivery. At the third day of life, tachypnoea and metabolic acidosis led to a detailed metabolic investigation. The investigation of amino acids in plasma revealed an increased concentration of glycine (722 \(\mu\)M; 80–440 \(\mu\)M). As for patient 1, urine organic acid analysis showed an increased concentration of \(\beta\)-hydroxybutyric acid (17818 mmol/mol of creatinine), \(\beta\)-hydroxypropionic acid (25096 mmol/mol of creatinine), \(\beta\)-hydroxyisovaleric acid (34 mmol/mol of creatinine), lactic acid (1322 mmol/mol of creatinine), propionylglycine, methylcitrate and tiglylglycine. Propionyl-CoA carboxylase activity in cultured skin fibroblasts was strongly decreased (0.3 nmol h\(^{-1}\) · mg of protein\(^{-1}\)), confirming the diagnosis of PA that was treated according to standard methods (see also the Patient 1 subsection).

The child always had a myopathic appearance and exercise intolerance. Standard exercise testing at age 5 years showed an increase in blood lactate from normal concentrations at rest to 20 mM lactate after a short run of approx. 50 m. This was the reason for further bioenergetic analyses in muscle tissue (Table 1).

**Table 1 Bioenergetic analysis in muscle tissue of PA patients**

<table>
<thead>
<tr>
<th>Experiment</th>
<th>Patient 1</th>
<th>Patient 2</th>
<th>Control range</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. Oxidation of pyruvate, malate and succinate</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>([1,4-14C])Pyruvate + malate</td>
<td>0.13</td>
<td>0.56</td>
<td>3.61–7.48</td>
</tr>
<tr>
<td>([1,4-14C])Pyruvate + carnitine</td>
<td>0.21</td>
<td>0.59</td>
<td>2.84–8.24</td>
</tr>
<tr>
<td>([1,4-14C])Malate + pyruvate + malonate</td>
<td>0.18</td>
<td>0.32</td>
<td>4.68–9.62</td>
</tr>
<tr>
<td>([1,4-14C])Malate + acetyl carnitine + malonate</td>
<td>0.46</td>
<td>1.32</td>
<td>3.43–7.30</td>
</tr>
<tr>
<td>([1,4-14C])Malate + acetyl carnitine + arsenite</td>
<td>0.43</td>
<td>1.45</td>
<td>2.05–6.39</td>
</tr>
<tr>
<td>([1,4-14C])Succinate + acetyl carnitine</td>
<td>0.27</td>
<td>0.82</td>
<td>2.54–6.39</td>
</tr>
<tr>
<td>B. Single enzyme activity</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Complex I</td>
<td>16</td>
<td>34</td>
<td>70–250</td>
</tr>
<tr>
<td>Complex II</td>
<td>59</td>
<td>38</td>
<td>67–177</td>
</tr>
<tr>
<td>Complex III</td>
<td>446</td>
<td>989</td>
<td>2200–6610</td>
</tr>
<tr>
<td>Complex IV</td>
<td>507</td>
<td>593</td>
<td>810–3120</td>
</tr>
<tr>
<td>Complex I + coenzyme Q + complex III</td>
<td>87</td>
<td>118</td>
<td>300–970</td>
</tr>
<tr>
<td>PDHc</td>
<td>6</td>
<td>10</td>
<td>34–122</td>
</tr>
<tr>
<td>PDHc-E1</td>
<td>1.08</td>
<td>1.52</td>
<td>1.23–4.13</td>
</tr>
<tr>
<td>PDHc-E3</td>
<td>353</td>
<td>533</td>
<td>760–2630</td>
</tr>
<tr>
<td>KGDHc</td>
<td>n.m.</td>
<td>3</td>
<td>23–49</td>
</tr>
<tr>
<td>C. ATP and phosphocreatine</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ATP + phosphocreatine</td>
<td>10</td>
<td>0.0</td>
<td>42–81</td>
</tr>
<tr>
<td>D. Mitochondrial reference enzyme</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Citrate synthase</td>
<td>81</td>
<td>147</td>
<td>37–162</td>
</tr>
</tbody>
</table>

Muscle biopsies

Muscle biopsies (musculus vastus lateralis dexter) and forearm skin punch biopsies were performed in both patients. Furthermore, fibroblast cultures from patients with confirmed diagnosis of PDHc (pyruvate dehydrogenase complex) deficiency (n = 3) and healthy volunteers (n = 12) were used. Biopsies and subsequent experiments were performed after receipt of informed consent. The study was approved by the Institutional Review Board (Medical Faculty of Heidelberg, Heidelberg, Germany). Further investigations were performed in muscle tissue (musculus quadriceps femoris) of wild-type C57Bl/6 mice. Animal care followed the official governmental guidelines and was approved by the government ethics committee.

**Human skin fibroblast cultures**

Human skin fibroblasts from forearm skin biopsies were cultivated under standard conditions in Dulbecco’s modified Eagle’s medium at 37°C supplemented with 10% (v/v) fetal calf serum, 100 \(\mu\)g/ml penicillin, 100 \(\mu\)g/ml streptomycin, 2.5 \(\mu\)g/ml of fungizone and 200 \(\mu\)M uridine until confluency [8]. Medium was changed twice a week. Each cell culture was tested for contamination with Mycoplasma spp. before enzyme analysis.

**Preparation of tissue extracts**

Fibroblast and muscle homogenates as well as SMPs (sub-mitochondrial particles) from bovine heart were prepared as previously described [8–10].

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PDHc activity

Spectrophotometric analysis of PDHc activity [E1 (pyruvate decarboxylase), EC 4.1.1.1; E2 (dihydrolipoate transacetylase), EC 2.3.1.12; E3 (dihydrolipoate dehydrogenase), EC 1.8.1.4] was performed in purified porcine PDHc (Sigma–Aldrich, Schnelldorf, Germany), in SMP, and in homogenates from human skin fibroblasts and quadriceps muscle biopsy specimens using a p-lodotnitrotetrazolium Violet-coupled system as previously described [8]. In brief, steady-state activities of PDHc were recorded using a computer tuneable spectrophotometer (SPECTRAMax Plus 340 microplate reader; Molecular Devices, Sunnyvale, CA, U.S.A.) operating in the dual wavelength mode. Samples were analysed in temperature-controlled 96-well plates in a final volume of 300 µl PDHc (purified enzyme, 160 mg units/ml; SMP, 1.2 mg of protein/ml; fibroblast homogenate, 4 mg of protein/ml) was assayed in a buffer containing 0.5 M potassium phosphate, 2.5 mM NAD+, 5 mM L-carnitine, 0.2 mM thiamine pyrophosphate, 0.1 M CoA, 0.1% (v/v) Triton X-100, 1 mM MgCl2, 1 mM EDTA, 0.5 M NADH, 6.5 mM phenazine methosulphate, which was adjusted to pH 7.5 at 30°C. PDHc activity was determined as p-lodotnitrotetrazolium Violet reduction at a wavelength of λ = 500–750 nm. The specificity of this assay was confirmed by complete inhibition of PDHc activity by the specific E1 inhibitor 3-fluoropyruvate (5 mM).

To investigate the effect of acetyl-CoA and propionyl-CoA, we varied the concentrations of these CoA esters (0–1 mM) and pyruvate (0–0.1 mM). Additional experiments were performed using short-chain (acetyl-CoA [C2], propionyl-CoA [C3] and butyryl-CoA [C4]), medium-chain (hexanoyl-CoA [C6], octanoyl-CoA [C8], and decanoyl-CoA [C10]) and long-chain (myristoyl-CoA [C14] and palmitoyl-CoA [C16]) acyl-CoA esters as well as corresponding short- and medium-chain fatty acids (up to 1 mM; all adjusted to pH 7.5).

Spectrophotometric analysis of KGDHc (α-ketoglutarate dehydrogenase complex) activity

Spectrophotometric analyses of KGDHc [KGDHc subunits: Elk (α-ketoglutarate dehydrogenase), EC 1.2.4.2; E2k (dihydrolipoil succinyltransferase), EC 2.3.1.61; and E3] were performed in SMP and muscle homogenates as previously described [10]. In brief, KGDHc (650 m-units/ml) was assayed in a buffer containing 35 mM potassium phosphate, 5 mM MgCl2, 0.5 mM EDTA, 0.5 mM NAD+, 0.2 mM thiamine pyrophosphate, 0.04 mM CoA-SH (where SH is thiol group) and 2 mM α-ketoglutarate, which was adjusted to pH 7.4 at 30°C. KGDHc activity was determined as NAD+ reduction at a wavelength of λ = 340–400 nm. The effect of propionyl-CoA on KGDHc activity was tested in analogy to PDHc (n = 8).

Spectrophotometric analysis of OXPHOS complexes I–V

The catalytic activities of respiratory chain complexes I–V in SMP and muscle homogenates were investigated as previously described [9–14]. The addition of standard respiratory chain inhibitors [complex I, 2-n-decylquinazolin-4-yl-amin (1 µM); complex II, thienoyltrifluoroacetone (8 mM); complex III, antimycin A (1 µM); complex IV, NaN3 (2 mM); complex V, oligomycin (80 µM)] revealed good inhibitory responses (93–100% of control activity, P < 0.001 versus controls), confirming a high specific enzyme activity in our assay system. Acyl-CoA esters (C2, C3, C4 and C6) and corresponding fatty acids (all adjusted to pH 7.4) were added and effects on single respiratory chain complexes I–V were determined at concentrations up to 1 mM each (complex I, 0.5 mg/ml of protein; complex II, 0.3 mg/ml of protein; complex III, 0.1 mg/ml of protein, complex IV, 0.01 mg/ml of protein; complex V, 1.1 mg/ml of protein). By analogy, complex III activity was also investigated in isolated complex III from bovine heart (0.1 mg/ml of protein).

Radiometric analysis of mitochondrial OXPHOS

Radiometric analysis was performed in freshly prepared quadriceps-muscle biopsy specimens from PA patients. Supernatants (600 µg) were prepared and oxidation rates of [1-14C]pyruvate, [U-14C]malate and [1,4-14C]succinate were determined as previously described [14–16].

ATP production

Spectrophotometric analysis of ATP production was determined in muscle biopsies of PA patients using unlabelled pyruvate, malate and succinate as described previously [14]. In brief, spectrophotometric analysis of ATP production is coupled with the formation of NADPH (λ = 320–400 nm, 25°C). The test principle consists of two enzyme reactions. In the first step, glucose (30.3 mM) and ATP are catalysed to glucose-6-phosphate and ADP by hexokinase (EC 2.7.1.1). Subsequently, glucose-6-phosphate dehydrogenase (EC 1.1.1.49) catalyses glucose-6-phosphate and NADP+ to 6-phosphogluconolactone and NADPH.

Protein electrophoresis of muscle tissue

One- and two-dimensional BN-PAGE (blue native PAGE) and subsequent in-gel activity assays and Western-blot analysis were performed in muscle biopsies of PA patients as described previously [17]. Antibodies against OXPHOS complex I (subunits 39 kDa and ND6 (NADH dehydrogenase 6)), complex II (subunit 70 kDa) and complex III (subunit Core 2) were obtained from Molecular Probes (Eugene, OR, U.S.A.), and against cyclophilin B, from Affinity Bioreagents (Golden, CO, U.S.A.).

Electron microscopy of muscle tissue

Electron microscopical examination of muscle biopsies of PA patients was done according to standard methods.

Data analysis

Enzyme activities were normalized to citrate synthase (EC 2.3.1.3) activity and the protein concentration of the same sample [18,19]. Results were expressed as means ± S.D. for at least three independent experiments. ANOVA followed by post hoc Bonferroni’s multiple comparison test (for three or more groups) or Student’s t test (for two groups) were used to calculate statistical differences between groups. Results are presented as the means ± S.D. if not indicated differently. Statistics were calculated using SPSS for Windows 12.0 software. P < 0.05 was considered significant.

RESULTS

Propionyl-CoA inhibits PDHc

In purified porcine PDHc, propionyl-CoA showed an inhibition of PDHc activity, being competitive with respect to pyruvate (Figure 1). Notably, this inhibitory effect was in the same range as for acetyl-CoA, the physiological product and known inhibitor of PDHc (Figure 2). For propionyl-CoA (0.125–1 mM), the inhibition constant (K) ranged from 101 to 422 µM [3–12 times K (Michaelis–Menten constant)] and for acetyl-CoA (0.125–1 mM), it ranged from 90 to 256 µM (2–7 times K). The K was 35 ± 0.5 µM for pyruvate. Similar results were found in SMP from bovine heart (results not shown).
Figure 1 Inhibition of PDHc by propionyl-CoA
Spectrophotometric analysis in purified PDHc from porcine heart demonstrates uncompetitive inhibition of PDHc activity by propionyl-CoA with respect to pyruvate (Pyr) as demonstrated by a Lineweaver–Burk plot. Values are given as the means ± S.D. (where no error bars are visible the S.D. is within the symbol). Experiments were performed at least in triplicate.

Figure 2 Inhibition of PDHc by acetyl-CoA
Spectrophotometric analysis in purified PDHc from porcine heart demonstrates uncompetitive inhibition of PDHc activity by acetyl-CoA with respect to pyruvate (Pyr) as demonstrated by a Lineweaver–Burk plot. Values are given as the means ± S.D. (where no error bars are visible the S.D. is within the symbol). Experiments were performed at least in triplicate.

Next, we determined PDHc activity in fibroblast homogenates from two PA patients. Assay conditions allowed enzyme analysis in the absence of intracellularly formed propionyl-CoA. Notably, fibroblasts from both patients revealed PDHc activities within the control range, confirming our hypothesis of secondary PDHc inhibition induced by propionyl-CoA (Figure 3). As expected, patients with confirmed PDHc deficiency showed a reduced PDHc activity in fibroblasts (Figure 3).

Acyl-CoA esters inhibit PDHc in a chain-length-specific manner
To investigate whether PDHc inhibition was dependent on the chain length of acyl-CoA esters, we investigated the inhibitory effect of similar monocarboxylic CoA esters ranging from C₄ to C₁₆ in purified porcine PDHc. In fact, PDHc was also effectively inhibited by C₄, C₆, C₈, and C₁₀ esters but less effectively by C₁₀ and C₁₆ esters (Figure 4A). In contrast, short-chain fatty acids (up to 1 mM) did not inhibit of PDHc (Figure 4B). These results were confirmed in SMP from bovine heart (results not shown).

DISCUSSION
The accumulation of metabolites from the alternative oxidative pathways of propionyl-CoA is the biochemical hallmark of PA.
Mitochondrial dysfunction in propionic aciduria

During metabolic crisis, additional biochemical abnormalities such as lactic acidosis, hyperketosis and hypoglycaemia indicate the development of severe mitochondrial dysfunction resulting in impairment of energy metabolism [1]. The present study significantly adds to the understanding of these mechanisms demonstrating (i) a severe bioenergetic disturbance and ultrastructural changes of mitochondria in muscle tissue of PA patients and (ii) propionyl-CoA-induced synergistic inhibition of PDHc, KGDHc and complex III in vitro. These results support the hypothesis that secondary mitochondrial dysfunction induced by accumulating toxic metabolites plays a major role in the pathomechanisms of PA [7]. We clearly demonstrated that propionyl-CoA, but not propionate, is the major toxic metabolite involved in this scenario.

PDHc is a bioenergetic bottleneck coupling cytosolic anaerobic glycolysis with the mitochondrial tricarboxylic acid cycle and OXPHOS. Not surprisingly, inherited PDHc deficiency results in a severe disturbance of energy metabolism and frequently in a fatal disease course [20]. In analogy, secondary inhibition of PDHc by propionyl-CoA can be suggested as an important mechanism inducing energy failure in PA, which is confirmed by severely reduced ATP and phosphocreatine concentrations in muscle tissue.
of PA patients and increased serum lactate concentrations. In line with this, multiple organ failure in PA patients predominantly manifests in tissues with a high energy demand, such as the central nervous system (encephalopathy, movement disorders and developmental retardation), skeletal muscle (myopathy), heart muscle (cardiomyopathy) and bone marrow (pancytopenia) [1,3]. Apart from mammalian species, propionyl-CoA-induced inhibition of PDHc has recently been suggested as a major mechanism underlying the antibacterial and antifungal properties of propionate [4]. Thus propionyl-CoA-induced mitochondrial dysfunction can be regarded as a common final pathway involved in different conditions, i.e. inherited disorders of propionate metabolism and propionate toxicity. Future studies should focus on concentration-dependent effects of propionyl-CoA in tissues of children affected with PA. However, since the amount of a muscle biopsy specimen in children is always very limited because of ethical and functional considerations, this important aspect could not be investigated additionally in the present study.

In addition to PDHc inhibition, alternative mechanisms for propionyl-CoA and metabolites of alternative propionate oxidation should be considered as relevant. In particular, inhibition of respiratory chain complex III (the present study) and the tricarboxylic acid cycle enzymes KGDHc (the present study), citrate synthase [5] and succinyl-CoA synthetase [4] may also contribute to impaired energy metabolism in this disease. Interestingly, inherited deficiency of succinyl-CoA synthetase caused by deleterious mutations in the SUCL2 gene has recently been associated with encephalomyopathy and mtDNA depletion [21], linking the tricarboxylic acid cycle with mtDNA homoeostasis [21,22]. In addition, increased oxidative stress, which has been demonstrated in an in vitro model for disorders of propionate metabolism, induces mtDNA damage [23,24]. Interestingly, the amount of mtDNA and the activities of OXPHOS complexes I, III and IV, which are partially encoded by mtDNA, were significantly decreased in muscle tissue of both PA patients; however, it remains unclear whether this result reflects a causal link. Besides mtDNA homoeostasis, other secondary or tertiary targets might be involved but have not yet been identified.

PA shares a variety of biochemical and clinical similarities with methylmalonic aciduria, which is caused by inherited deficiency of methylmalonyl-CoA mutase or the synthesis or transport of its cofactor, 5’-adenosylcobalamin [1]. We have recently hypothesized that propionyl-CoA and metabolites deriving from propionyl-CoA, such as 2-methylcitrate, might act as endogenous neurotoxins also in this disease, whereas methylmalonate most likely plays a minor role [13,14,25]. Since the manifestation of secondary metabolic blocks is pathophysiological relevant in PA and methylmalonic aciduria, it is of interest to investigate whether alternative energy substrates such as succinate and citrate might be beneficial for metabolic maintenance treatment and intensified emergency treatment of these patients helping to restore mitochondrial energy metabolism and to prevent multiple organ failure.

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