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**Pyruvate carboxylase deficiency type A and type C: characterization of 5 novel pathogenic variants in PC and analysis of the genotype-phenotype correlation**

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This article has been accepted for publication and undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1002/humu.23742.

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Abstract

Pyruvate carboxylase deficiency (PCD) is caused by bi-allelic mutations of the PC gene. The reported clinical spectrum includes a neonatal form with early death (type B), an infantile fatal form (type A), and an late onset form with isolated mild intellectual delay (type C). Apart from homozygous stop-codon mutations leading to type B PCD, a genotype-phenotype correlation has not otherwise been discernible. Indeed, patients harboring bi-allelic heterozygous variants leading to PC activity near zero can present either with a fatal infantile type A or with a benign late onset type C form.

In this study, we analyzed six novel patients with type A (three) and type C (three) PCD, and compared them with previously reported cases. Firstly, we observed that type C PCD is not associated to homozygous variants in PC. In-silico modeling was used to map former and novel variants associated to type A and C PCD, and to predict their potential effects on the enzyme structure and function. We found that variants lead to type A or type C phenotype based on the destabilization between the two major enzyme conformers. In general, our study on novel and previously reported patients improves the overall understanding on type A and C PCD.

KEY WORDS

Pyruvate carboxylase deficiency; Neuro-developmental delay; Pyruvate carboxylase; Biotin carboxylase domain; Carboxyl transferase domain; PC tetramerization domain; Biotin carboxyl carrier protein domain

1. INTRODUCTION

Pyruvate carboxylase deficiency (PCD) (MIM# 266150) is a rare recessive genetic disorder (estimated prevalence 1:250.000) affecting the confluence of glucose metabolism, lipogenesis and neurotransmitter synthesis (Wang et al., 2009). The
pyruvate carboxylase (PC) gene maps to 11q13.2 and consists of 4 non-coding exons and 20 coding exons. No pseudo-genes are known. Three tissue-specific variant transcripts differ from each other at the 5'-UTR region but encode the same 1178 amino acid protein (in addition to the 20 amino acid mitochondrial leader sequence) containing 3 domains (biotin carboxylation, BC; transcarboxylation, CT; biotin carboxyl carrier, BCCP) and 2 spacer regions (Wallace et al., 1998). Pyruvate carboxylase (PC) is a homotetramer localized to the mitochondrial matrix and converts pyruvate into oxaloacetate. PC structures resolved by means of X-ray crystallography suggest that the protein undergoes large conformational transitions over the course of its two-step enzymatic reaction (Maurice et al., 2007; Xiang et al., 2008). In the first step, a biotin moiety covalently bound to the BCCP domain is carboxylated. This step requires ATP, which binds to the BC domain and is allosterically activated by acetyl-CoA. In the second step, biotin transfers a carboxyl group to pyruvate. This reaction takes place at the CT domain. The active sites of the BC and CT domains are separated by more than 70 Å in the PC structures of Staphylococcus Aureus and Rhizobium Etli. Biotin translocation requires large scale molecular motion of the BCCP domain between the BC and CT domains.

Mitochondrial oxaloacetate participates in several critical biochemical pathways. It is converted into aspartate, required for protein synthesis, for the production of argininosuccinate, a metabolite of urea cycle, and for the conversion of glutamate into the neurotransmitter γ-aminobutyric acid (GABA). Oxaloacetate is a component of the tricarboxylic acid (TCA; Kreb) cycle, and its absence leads to a reduction of mitochondrial NADH, a substrate of complex I of the respiratory chain. During fasting, oxaloacetate is converted into phosphoenolpyruvate, required for gluconeogenesis. Finally, oxaloacetate is necessary for the transport of mitochondrial acetyl-CoA into the cytoplasm, where it participates in the synthesis of fatty acids. Taken together, lack of
oxaloacetate due to PCD has a plethora of severe metabolic consequences (Marin-Valencia et al., 2005).

Two clinical subtypes with early onset and fatal outcomes have been defined: A (also called ‘American type’) and B (also called ‘French type’). Type A PCD manifests several months after birth and presents with increasing generalized developmental delay and hypotonia, failure to thrive, profound apathy and intermittent lactic acidemia with intercurrent illnesses or other physiologic stress (e.g. infections). All reported patients have died during infancy or early childhood. Few patients with this infantile and always fatal PCD form were reported in the literature (Robinson et al., 1987; Ahmad et al., 1999; Wang et al., 2008; Monnot et al., 2009).

Type B PCD presents in the first hours after birth with severe lactic acidosis, hyperammonia, hypercitrullinemia and hypoglycemia. Immediately following birth, neurological status and consciousness level are typically normal, but after 1 to 72 hours a hypokinetic-rigid syndrome with profound axial hypotonia appears, which is accompanied by tremor of the limbs and bizarre ocular movements in some patients. During the first months of life, increasing liver insufficiency and seizures develop in some patients. Patients die within five months after birth (Carbone et al., 1998; Carbone et al., 2002; Ostergaard et al., 2012). Only 38 inactivating mutations in the PC gene have been described in patients with type A and B PCD (Carbone et al., 1998; Carbone et al., 2002; Wang et al., 2008; Monnot et al., 2009; Ostergaard et al., 2012; Ortez et al., 2013; Breen et al., 2014; Wexler et al., 1998). Several other PCD patients have been reported based on biochemical stigmata of PCD and/or on reduction of pyruvate carboxylase activity, but without molecular confirmation of the diagnosis (Oizumi et al., 1983;
Therapeutic options for type A and B disease are limited. Anaplerotic diet or compounds (e.g. triheptanoin, citrate, or aspartate) have shown variable success in reducing metabolic derangements, but have not achieved an amelioration of the neurological symptoms and a prolongation of life span (Garcia-Cazorla et al., 2005; Breen C et al., 2013; Mochel et al., 2005).

A third form of PCD (type C or benign) is characterized by short episodes of ketoacidosis during infancy and childhood, typically associated with intercurrent illnesses, mild psychological delay and neurologic symptoms, and survival until late adolescence in the six reported patients (Van Coster et al., 1991; Higgins et al., 1994; Higgins et al., 1997; Hamilton et al., 1997; Arnold et al., 2001; Schiff et al., 2006; Wang et al., 2008). PCD mutations were reported in only two cases (Wang et al., 2008). Enzymatic activity was reported as 1-10% (of healthy control) in skin fibroblasts or peripheral blood leukocytes in three patients (Van Coster et al., 1991; Hamilton et al., 1997; Arnold et al., 2001), below 1% in two patients (Schiff et al., 2006; Wang et al., 2008) and was unreported in one patient (Wang et al., 2008). Thus, residual enzymatic activity does not appear to be a reliable prognostic parameter in this condition.

Reported genotype/phenotype correlations in both fatal PCD forms (infantile type A and neonatal type B) have been inconsistent. Initially it appeared that the presence of at least one truncating mutation in the PC gene led to type B presentation, while bi-allelic missense mutations lead to type A (Carbone et al., 1998; Carbone et al., 2002; Monnot et al., 2009). However, additional reports have not been as clear, with type B presentation seen in patients with bi-allelic missense mutations (Wang et al., 2008; Ostergard et al., 2008).
Moreover, Wang and colleagues hypothesized that mosaicism for mutations in CNS tissues correlated with patient survival (Wang et al., 2008).

The genetic difference between type A and C PCD has not been elucidated. Although bi-allelic mutations in $PC$ have been reported in both, there has not been a clear correlation of genotype with phenotype. The prognosis among type A and type C patients is generally different, dying the former within the first years of life and surviving the latter sometimes until adulthood.

In the current study, we analyzed clinical, electrophysiological, neuroradiological, biochemical, and genetic findings in six patients with PCD, three with type A (patients 1, 2, 3) and three with type C (patients 4, 5, 6) PCD. The patients harbor five novel and one known pathogenic variant in $PC$. We compared the genetic features of these six novel patients with the previously reported type A and type C patients. We created homology models for the full structure of human PC representing two main conformations of the enzyme, where either the BCCP domain is in the vicinity of BC domain or the BCCP domain is adjacent to the CT domain, and we mapped most of the reported mutations associated with type A and type C PCD onto the models. Together with interpolation of motion between the two structural states of the enzyme, the mapped mutations provided structural insights into the potential effects on the enzyme's function.

Taken together, this study aims to shed further light into the still unclear genotype - phenotype correlation for type A and type C PCD and to provide future PCD patients with a clearer prognosis basing on the revealed genetic mutations.
2. PATIENTS AND METHODS

2.1 Patients
Six patients were recruited worldwide from treating metabolic medicine physicians and were included in the study according to clinical and biochemical signs of type A or type C PCD.

2.2 MRI scan
For patients 1, 3 and 4, brain MRI was performed on a 3 Tesla MR scanner using a standard clinical protocol. For patient 2, brain imaging with spectrography was performed on a 1.5 Tesla MR scanner using a standard clinical protocol.

2.3 Gene sequencing

2.4 PC enzymatic activity
In patients 2, 3 and 4, the enzymatic activity of pyruvate carboxylase activity was measured in cultured skin fibroblasts from patients 2, 3 and 4 according to standard protocols [Robinson et al., 1985; Atkin et al., 1979]. In patients 1, 5 and 6, the enzymatic

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activity in skin fibroblasts could not be measured due to either technical issues or lack of parental consent.

2.5 Pyruvate carboxylase modeling and variant mapping
Two major conformers of the PC enzyme were modeled to mimic the functional states of biotin carboxylation (conformer 1, PDB ID 2qfc (Maurice et al., 2007)) and carboxyl transfer to pyruvate (conformer 2, PDB ID 3bg3 and 3bg5 (Xiang et al., 2008)). In the conformer 1, BCCP domain is adjacent to the BC domain; in the conformer 2, BCCP domain is adjacent to the CT domain. All the homology modeling was performed using Modeller 9.15 (Sali et al., 1993). The previously reported mutations as well as the novel mutations associated with type A and type C PCD were mapped onto the modeled structures. The structures were visualized by means of Pymol (Schrödinger). The artificial motion connecting the two modeled conformers was constructed by linearly interpolating between the structures along the difference vector between them. For a detailed description of PC modeling methods, we refer to “supplementary material and methods”.

2.6 Free energy calculations and PC stability
The changes in PC stability upon each mutation were estimated by the established Rosetta protocol (Kellogg et al., 2011) using the Cartesian version of the method (Alford et al., 2017). Two Rosetta energy functions were used for calculations (Talaris2014 and ref2015 (Alford et al., 2017)): the calculated values are reported as a range between the two obtained estimates. The free energy difference (ΔΔG) was calculated upon each type A and type C variant previously reported in the literature as well as in this study. The ΔΔG values were separately calculated for both conformers 1 and 2, including all the possible monomer combinations for the heterozygous and homozygous variant scenarios.
The difference (ΔΔΔG) among ΔΔG_{conformer1} and ΔΔG_{conformer2} is used as an indicator of the PC stability and indirectly of its functional impairment. It can be assumed that evolutionary constraints have optimized the relative stability between the conformers for carrying out the enzyme’s function and a distortion of this balance, be it destabilization or over-stabilization of one conformer with respect to the other, may be viewed as an indication of the functional impairment of pyruvate carboxylase. Positive ΔΔΔG values indicate a relative stabilization of the conformer 2 with respect to the conformer 1 and negative ΔΔΔG values indicate on the contrary that the conformer 1 is more stable than the conformer 2. For a detailed description of the mathematical basis of the used thermodynamic model and the Rosetta based calculations we refer to “supplementary material and methods”.

3. RESULTS

3.1 Clinical, biochemical, electrophysiological and neuroradiological features of the patients

Patient 1 (male) is the second child of consanguineous parents and brother of a 2 years old healthy boy. Delivery at 37 weeks of pregnancy was uncomplicated, and birth weight, length and head circumference were within the normal limits. No problems were noted until 5 months of age, when he was admitted to hospital due to an acute bronchopneumonia with severe tachyypnea, fever and somnolence. He had severe lactic acidosis (serum lactate 18.1 mmol/l) and hyperglycemia (136 mg/dl). Ammonia was 53 μmol/l (normal range 30-60). Organic acids analysis in urine identified elevated lactate and ketones. Initially a defect in pyruvate dehydroxygenase complex was suspected. The patient was treated with intravenous bicarbonate and he was put on a high fat diet (50-69%). Acidosis improved but was never completely corrected, with elevated serum lactate (7.6 mmol/l) and base excess (-4.3 mmol/l). At 7 months of age, acute
metabolic acidosis occurred with RSV bronchiolitis (serum lactate 9.7 mmol/l). Currently, the patient is 11 months old with weight of 8280 g (10th percentile), length 71 cm (3rd percentile), and head circumference 45 cm (3rd percentile). His developmental milestones correspond to 5-months of age and he is slightly dystonic. Brain MRI performed at 6 months showed an increased signal in putamen, globus pallidus and caudate bilaterally (Figure 1).

Patient 2 (female) is the first child of non-consanguineous Aboriginal Canadian parents. At birth, she suffered from lactic acidosis and generalized hypotonia. Birth weight and length were within the normal range, while head circumference was 39 cm (3 cm above 97th percentile). Until the fifth year of age, she was admitted four times due to acute metabolic acidosis and dehydration during minor intercurrent illness. Metabolic studies revealed continuously elevated levels of serum lactate (5 - 7.5 mmol/l and maximal peak 15 mmol/l during an infection), serum pyruvate (maximal peak 0.28 mmol/l) and alanine (750 - 1500 µmol/l). Base excess and pH were within the normal range between illnesses and decreased during infection-associated metabolic misbalances, the lowest measured base excess was -13 and the lowest pH was 7.13. At 2 days of age, brain MRI revealed ventriculomegaly with large subependymal cysts and pronounced serum lactate peaks in spectroscopy. Currently, she is 11 ½ years old, has a weight of 45.9 kg (75th percentile), length of 147.5 cm (50th percentile) and head circumference of 52.5 cm (50th percentile). Developmental progression has arrested at approximately the stage of a 2-years old child. The patient is able to pronounce about 10 words, but is unable to build 2-words sentences. She can walk and run with good coordination, and does have problems with tantrums. No psychological testing has been performed and no further developmental progression is expected. The patient has been treated with oral biotin, aspartate, carnitine, thiamine and citrate since the diagnosis.

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Patient 3 (male) is the first child of non-consanguineous parents. The birth was uncomplicated at the 39th pregnancy week. At 3 weeks of age, bilateral clonic seizures started, lasting from 30 seconds to 3 minutes and sometimes occurring in clusters of 3-4 seizures. Oral Levetiracetam and Vigabatrin partially controlled the seizures; oral steroids were added for 30 days due to breakthrough infantile spasms. At 6 months of age, he suffered two episodes of metabolic acidosis within a few weeks without apparent trigger. He was somnolent and hypotonic, had increased serum lactate (10.8 mmol/l), decreased bicarbonate (11 mmol/l) and pH (7.35), normal ammonia and glucose levels and was discharged after intra-venous fluid therapy. Plasma alanine, taurine, glutamic acid, glutamine and proline were almost constantly elevated, while aspartic acid, threonine, citrulline, cystine, lysine and histidine were intermittently elevated. In the unique CSF test, alanine and citrulline were elevated. MRI investigation at 2 months of age revealed cystic change in the periventricular white matter, delayed myelination and underopercularization of the sylvian fissures (Figure 1). At the current age of 22 months, the patient fixes objects but does not actively follow them, brings the hands to midline, takes objects with both hands and cannot sit without support. Weight and head circumference are on the 50th percentile and length is on the 75th percentile. He is currently treated with oral Biotin (10 mg daily).

Patient 4 (male) is the first child of non-consanguineous parents. Fetal MRI revealed ventriculomegaly with septations and agenesis of the corpus callosum (Figure 1). Birth at the 40th pregnancy week and neonatal period were uncomplicated, apart from hyperbilirubinemia. At 11 months, he suffered from an acute ear infection, and then abruptly developed mental status changes, ultimately becoming comatose, and Kussmaul breathing. Having a pH 6.7 and bicarbonate 2 mmol/l, the patient was resuscitated with i.v. bicarbonate infusion. After this first metabolic crisis, he had permanently elevated
levels of urinary ketone bodies and plasma pyruvate and lactate. Serial biochemical tests revealed constantly increased levels of plasma alanine and intermittently elevated levels of plasma proline and lysine. At 3, 4 and 5 years of age, the patient suffered from acute metabolic acidosis episodes in the setting of cough, RSV infection and gastroenteritis, respectively. At the current age of 6 years, he has stereotyped movements (hand flapping), walks, jumps, climbs stairs, throws and catches. He expresses about 10 purposeless sounds but not intelligible words, follows commands and seems to understand speech, plays and shares with other children, distinguishes strangers from known people. Weight and length are on the 90th percentile. The patient is currently treated with biotin (10 mg daily), carnitine (50 mg/kg daily) and riboflavin (50 mg daily).

Patient 5 (male) is the second child of non-consanguineous parents, born to non-consanguineous parents. He was born at full-term with normal weight and length. Head circumference was at the 99th percentile. Family’s clinical history was unremarkable. At 2 years of age, the child had good motor skills though he walked on his toes, but spoke only a few words. He was given a diagnosis of autism spectrum disorder because of poor eye contact and obsessive behaviour. He subsequently presented at that age to the emergency room with increased work of breathing and significant retractions with afebrile illness and upper respiratory infection. Metabolic evaluation revealed pH 7.07, bicarbonate 6 mmol/l, serum lactate 8.27 mmol/l, and ammonia 45 µmol/l which normalized with therapy. After paediatric intensive care with intravenous bicarbonate and fluid therapy, pH stabilized to normal. Retrospectively, a smell of ketosis on his breath was reported. He subsequently has had two additional episodes of metabolic acidosis with increased serum lactate (13 and 16 mmol/l) corrected with intensive fluid therapy. Currently, he is 3 years old, has a weight of 15.5 kg (67th percentile), length of 99 cm (72th percentile), and head circumference of 52 cm (98th percentile). After starting
oral triheptanoin (30 grams daily, ~2 g/kg/d) at 3 years of age, he is reported to have increased speech and improved eye contact.

Patient 6 (male) is the younger brother of patient 5. He was born full-term without complication and has not had any motor delay. Because of his brother’s clinical history, he was tested at birth and found to have the same PC variants as his older brother after his diagnosis. He is currently 16 months old, and his weight is 10.8 kg (60th percentile), length 77.5 cm (17th percentile), and head circumference 47.7 cm (71st percentile). He is babbling and interacting more than his brother, however he is not yet pointing. He has not had any episodes of metabolic decompensation even during illnesses. On routine testing, the patient 6 has increased lactic acid (6.3 mmol/l) and decreased bicarbonate (17 mmol/l). He was started oral triheptanoin at age 2 months and is currently getting 15 mg/d (1.5 g/kg/d).

3.2 PC variants and enzymatic activity of pyruvate carboxylase

In patient 1 (type A), PC harbors the duplication of c.449_451GGA in homozygous state. Both parents are heterozygous for c.449_451GGA. The encoded protein includes an additional glycine 151 after wild-type glycine 150. Enzymatic activity of pyruvate carboxylase could not be measured due to technical issues of testing.

Genetic testing of patient 2 (type A) revealed the homozygous variant c.1828G>A in PC, which causes the exchange of alanine 610 into threonine. Both parents are heterozygous for this variant. This mutation was already described in Indo-Americans from Canadian Cree and Ojibway tribes and was only associated to type A PCD. Enzymatic activity of pyruvate carboxylase measured in skin fibroblasts was 0.02 nm/min/mg protein (2.3% of control sample; control = 0.87+/−0.32 nm/min/mg) (Figure 2).

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Patient 3 (type A) harbors c.1877G>A in one PC allele; the other variant could not be identified within the coding sequence and the boundary regions of the other allele and was suspected to be deep-intronic. The c.1877G>A variant was paternally inherited and causes the exchange of arginine 626 into glutamine. Enzymatic activity of pyruvate carboxylase measured in skin fibroblasts was 0.09 nm/min/mg protein (6% of control sample; control = 1.5 nm/min/mg) and PC/CS ratio was 0.18 (4% of control sample) (Figure 2).

Patient 4 (type C) harbors c.797C>A in one PC allele. This variant causes the exchange of serine 266 into tyrosine and was maternally inherited. In patient 4 too, the second variant could not be identified neither in the coding part nor in the boundary regions of PC and was suspected to be deep-intronic. Enzymatic activity measured in skin fibroblasts was 0.00 nm/min/mg protein (0% of control sample) and PC/CS ratio was 0 (0% of control sample) (Figure 2).

The sibling patients 5 and 6 (type C) harbor two compound-heterozygous variants in PC, c.3242A>G of maternal origin and c.751+4A>G of paternal origin. The first variant causes the exchange of asparagine 1081 into serine and the second variant is likely expected to impair the splicing of intron 5 and the synthesis of the correct polypeptide. Both variants were previously described neither as pathogenic nor as benign variants. Enzymatic activity of pyruvate carboxylase could not be measured due to missing compliance of patients’ parents (Table 1).

### 3.3 Structural modeling of human PC and variant mapping

The constructed homology models represent two conformers of the homotetramer: conformer 1 with the BCCP domain adjacent to the BC domain and conformer 2 where BCCP brings biotin to the CT domain. The two conformers themselves contain several
sub-states. In the conformer 1, monomers A and B capture the position of BCCP domain between BC and CT domains (Figure 3) and monomers C and D have the BCCP domain closer to the BC domain (Supp. Figure S1): this state could be visited just before or right after carboxyl transfer to biotin. Conformer 2 also represents a combination of two sub-states. In all four monomers (A, B, C, D) of the conformer 2, the BCCP domain has biotin delivered to the CT domain. However, the conformation of the BCCP domain differs between the monomers A, B (Figure 3) and monomers C, D (Supp. Figure S1).

The overall motion of the pyruvate carboxylase performed during the carboxyl transfer step can be viewed in the supplementary section (Supp. Video S1). The movement of the enzyme covers more than 6.4 nm root mean squared displacement, when considering only the amino acid backbone atoms.

The overall variant mapping does not allow a unique classification of the phenotypic manifestation based on the localization of the variant (Figure 3). The variants associated with type A PCD appear in the BC and CT domains, whereas type C variants are found in the BC, CT and PT domains. Some variants are found near the ligand binding sites, e.g. p.G150dup (type A) and p.S266A (type C) close to ATP, p.R451C (type A) close to acetyl-CoA, p.A610T (type A) close to pyruvate and biotin, p.M743I (type A) close to pyruvate. Other variants are located at the interface of the enzyme’s monomeric subunits, e.g. p.R62C (type A), p.R451C (type A) and p.N1081S (type C) are in the proximity of residues in an adjacent monomer.

### 3.4 Predicted PC protein stability

In order to estimate how the single variant changes the PC stability and whether the protein stability degree is related to the disease phenotype, for each variant we have calculated the total double free energy difference ($\Delta\Delta G$), double free energy difference
of conformer 1 ($\Delta \Delta G_{\text{conformer1}}$), double free energy difference of conformer 2 ($\Delta \Delta G_{\text{conformer2}}$) and the difference between the latter two ones ($\Delta \Delta \Delta G$) as a relative stability indicator between the two conformers (Table 2). The calculations were performed using two energy functions yielding a range of stability estimates which in turn provide an indication for prediction accuracy. In the two following sections, we report the calculated free energies for all previously reported as well as novel variants: p.R62C, p.V145A, p.R156Q, p.R270W, p.R451C, p.A610T, p.R626Q, p.R631Q, p.M743I, p.A847V (all type A variants) and p.S266A, p.S266Y, p.T569A, p.N1081S (all type C variants).

**Variants in Type A Patients**

The free energy calculation suggests that the p.R62C variant not only does not distort the protein’s fold, but may even have a stabilizing effect. However, a change between the conformers is introduced by the p.R62C variant, resulting in $\Delta \Delta \Delta G$ of -3.1 to -1.0 kcal/mol and favoring the BCCP domain interaction with the BC domain. The p.R62C variant has been referred in association to type A PCD when found in a mosaic compound-heterozygosity with p.R631Q and p.A847V variants (Table 1). p.R631Q ($\Delta G$ 2.5 kcal/mol) has a low destabilizing effect on the enzyme, while p.A847V affects the relative monomer stability ($\Delta \Delta G$ 1.2 - 1.5 kcal/mol) suggesting their detrimental influence on the $PC$ function, which could be worsened by the p.R62C mutation.

The p.R631Q variant has also been implicated in another type A PCD case in compound heterozygosity with the p.R270W variant. While p.R631Q, as mentioned above, has a low destabilizing effect, p.R270W shows a potential difference among the two conformers ($\Delta \Delta \Delta G$ 0.0 - 6.6 kcal/mol), which may occur due to the stabilization of conformer 2 (BCCP domain interacting with the CT domain) (Table 2). The variant
destabilizes the interaction between BCCP and BC domains (conformer 1), which could perturb the balance between the enzyme conformations.

p.R156Q in the BC domain has a destabilizing effect on the protein, as well as altering the balance between the conformers. p.R156 is close to the ATP binding site.

The homozygous p.M743I variant has a strong destabilizing effect (ΔΔG 11.0 - 29.6 kcal/mol). The destabilization is prominent in both conformers.

The free energy calculations suggest that the p.A610T mutation is destabilizing (ΔΔG 2.3 - 6.7 kcal/mol). The major contribution to the decrease in PC stability comes from conformer 1, where biotin is interacting with the BC domain and is not in direct contact with the variant site. This suggests that this variant may interfere with the pyruvate-enzyme interaction.

The impact of p.R626Q variant to type A PCD is difficult to explain by the calculated changes in protein stability as the predictions based on the different energy functions differ substantially. This variant appears in compound heterozygosity with an intronic mutation and it is not excluded that PCD is also caused by an incorrectly spliced protein encoded from the second allele.

p.V145A variant is strongly destabilizing as predicted by the free energy calculations (ΔΔG 17.0 - 26.1 kcal/mol). It also substantially alters the balance between the preferred conformational states in comparison to the WT enzyme: conformer 2 appears to be more stable. The p.V145 residue is in the BC domain, however it is not in direct contact with biotin.

p.R451C variant is adjacent to the acetyl-CoA binding site. The current calculations do not provide an unambiguous prediction on the mutation’s effect on the enzyme’s stability.
stability, nor on the balance between the conformers. As a word of caution, it has to be
remarked that for the structural modeling of acetyl-CoA molecule we used the partial
structure of the non-hydrolyzable analog ethyl-CoA resolved in the pyruvate carboxylase
from *Rhizobium etli* (PDB 2QF7). This approximate representation of the ligand may not
warrant accurate estimation of the free energy changes.

In the computational study, we could not estimate the effect of G150 duplication on *PC*
stability, since modeling amino acid duplication would require to perform downstream
sequence shifting. The residue G150 is located in the BC domain adjacent to the ATP
binding site.

*Variants in Type C Patients*

Variant p.S266Y destabilizes the protein (ΔΔG of 3.4), while it was not possible to
obtain a reliable prediction of the effect of p.S266A on the enzyme’s stability. Neither of
the variants affects the balance between the conformers in the heterozygotic state (ΔΔΔG
0 kcal/mol and ΔΔΔG -2.2 - 0.2 kcal/mol, respectively). S266 residue is located in the
BC domain. However, the serine side-chain is facing away from the ATP and biotin
binding sites: therefore, the contribution of both mutations is mainly due to the altered
interactions with the neighboring residues in the BC domain (Figure 3 and Supp. Figure
S1).

Residue p.T569 is located in the CT domain and is not in direct contact to any ligand
important for the PC function. The mutation p.T569A is destabilizing (ΔΔG 3.3 - 3.4
cal/mol) and does not affect the balance between conformer 1 and conformer 2 (ΔΔΔG
0.0 kcal/mol).

Similarly, variant p.N1081S has a small destabilizing effect on the protein (ΔΔG 3.3 - 3.4
cal/mol). N1081 is located in the PT domain and is in direct contact with the interface
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residues of another PC monomer (Figure 3 and Supp. Figure S1), therefore it may play a role in the PC tetramerization. Interestingly, in one of the modeled states N1081 is placed closely to the R62 residue of another monomer (Supp. Figure S1) and, as discussed in the previous section, p.R62C variant has an effect on inducing type A PCD. p.N1081S variant may also have an effect on the acetyl-CoA binding: depending on the state considered, the distance between N1081 and the partial structure of ethyl-CoA is smaller than 0.5 nm.

Taken together, three out of four amino acid exchanges harbored by all reported type C patients have an overall small destabilizing effect to the enzyme (for one case, the effect on stability could not be reliably determined) and do not affect the balance between the major PC conformers. The amino acid exchanges present in type A patients manifest in a broad spectrum of the protein stability changes. The type A mutations in most cases appear to substantially disrupt the balance between the major conformational states of the enzyme, indicating a potentially detrimental effect of the genetic variant.

4. DISCUSSION

Pyruvate carboxylase deficiency is a rare neuro-metabolic disease with poor correlation of genotype to phenotype, leading to difficulty in predicting prognosis for patients. A previous study suggested that variants in the 1/3 C-terminal end of the protein could be tolerated and were unlikely to lead to disease since pathogenic missense mutations had been identified only in the 2/3 N-terminal end of the protein (Monnot et al., 2009). Additional reports have countered this observation, with compound-heterozygous mutations (p.A847V and p.L1137VfsX1170) in two patients with type A and one with type C PCD (Wang et al., 2008). We identified an N1081S variant in two patients with type C disease. Since all PCD reports from the pre-genome era (Oizumi et al., 1983;
Pineda et al., 1995) as well as several others from the genome era (Van Coster et al., 1998; Brun et al., 1999; Garcia-Cazorla et al., 2005) did not describe the pathogenic mutations underlining the phenotypes, any robust hypothesis about genotype-phenotype correlation was difficult.

Some genotype-phenotype correlations seem to be valid. The mutation c.1828G>A (p.A610T) appears to lead to type A disease, and all thirteen Indo-American patients from the Canadian Cree and Ojibway tribes with this variant had infantile onset of symptoms and death within the first 5 years of life (Carbone et al., 1998; Carbone et al., 2003). We reported a further Canadian patient with this mutation and type A phenotype.

p.A R631Q mutation has twice been described in patients with type A disease (Monnot et al., 2008; Wang et al., 2009). However, since this variant was in compound heterozygosity with p.R270W in one patient and with p.R62C and p.A847V in another patient, definitive conclusion about of the p.R631Q variant remain speculative. Apart from the p.A610T variant in Canadian First Nation people, no other founder variants have been described in other populations.

As stated above, only six patients with the milder late-onset type C PCD were reported before our study and the PCD causing variants were described in only two of them (Van Coster et al., 1991; Hamilton et al., 1997; Arnold et al., 2001; Schiff et al., 2006; Wang et al., 2008). Thus, our three type C patients significantly increase our ability to examine mutations in these mild patients. Patient 4 was found to have a heterozygous mutation of serine 266 (p.S266Y), a residue previously reported to be altered in a type C patient (p.S266A) (Wang et al., 2009). In our patient serine is substituted by tyrosine (p.S266Y) and in the previously reported patient serine was substituted by alanine (p.S266A), two very different amino acids. Serine 266 is the only amino acid, whose exchange has been
reported more than once in association with type C disease. The difference between the two variant amino acids together with the mild phenotype in both patients suggests that the exchange of this residue only mildly alters PC function and causes a mild phenotype. This hypothesis is supported by the fact that the second variant allele in the previously reported type C patient leads to a stop-codon (p.S705X), thus leading to only one allele in this patient capable of making an abortive protein (Wang et al., 2009). Although two patients are not enough to draw any definitive conclusion, they give us an initial hint on the functional consequences of serine 266 exchange. No other mutations have been reported in more than one type C patient, and thus conclusions about their pathogenicity and severity cannot be made. Regarding patients 5 and 6, at the current state it is difficult to speculate about the dysfunctional role of the variant c.3242A>G (p.N1081S) just considering its position, since exchange of N1081 residue has never been reported before.

Four of six patients (patients 3, 4, 5 and 6) are compound heterozygotes, with one missense variant and one intronic variant in PC. In patients 5 and 6, the intronic variant maps at the donor splice site of intron 5. In patients 3 and 4, the intronic variant has not been localized and was estimated to have a deep-intronic position. One of these patients (patient 3) has the type A phenotype, while the rest (patients 4, 5, 6) have the type C phenotype. Reviewing all five (former two and novel three) type C patients with revealed variants in PC, we observe that none of them harbors homozygous variants, rather all have compound-heterozygous variants. A hypothesis explaining this phenomenon might suggest that the polypeptide harboring one of the two compound-heterozygous variant amino acids is able to form a functional homotetramer, which is able in turn to exploit a sufficient enzymatic level and to produce a milder (type C) phenotype.

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However, it remains unclear why some patients with compound-heterozygous variants present the fatal type A phenotype. Possibly, both compound-heterozygous variants harbored by these type A patients lead to highly impaired monomers and therefore to an overall enzyme dysfunction. To shed light on this patho-physiological hypothesis and more generally on the phenotypical consequence of homozygous missense variants in \( PC \), it would be helpful to study patients with homozygous exchanges of residue S266 or residue T569. In fact, the computed free energy changes for these two type C associated mutations (p.S266Y and p.T569A), assuming them to be in a homozygous state, suggest a strong destabilization of the protein (more than 10 kcal/mol for each variant and conformer) (data not shown). Furthermore, for the homozygous mutations p.S266Y and p.N1081S the balance between the monomers is predicted to be disturbed as well (data not shown). The observation of patients with homozygous variants of S266 and T569 residues - associated in heterozygous state to type C phenotype in our study - would reveal whether a severe type A phenotype rises up, either caused by the overall destabilization of the enzyme or by the disturbed balance between the major conformational states of PC or by both effects.

Another interesting scenario that could provide insight into the difference between the type A and type C phenotypes, would be an identification of the p.R631Q mutation in a homozygous state not coupled with any additional mutations. Looking at the calculated free energy differences, p.R631Q appears to be only mildly destabilizing and to not perturb the balance between the major \( PC \) conformers. This observation would suggest its classification as a type C variant. However, coupling of this mutation to p.A847V or p.R270W results in a strong \( PC \) destabilization or in a disturbed balance between the conformers and finally in a type A phenotype (Monnot et al, 2009; Wang et al., 2008).
Independently from the associated phenotype, the mutated amino acids from patients 2, 3, 4, 5, 6 are highly conserved throughout the evolution (Supp. Figure S2).

Several reports have concluded that there is no correlation between residual enzymatic activity in fibroblasts and patient phenotype (Stern et al. 1995). We were only able to measure fibroblast pyruvate carboxylase activity in patients 2 and 3 (type A phenotype) and 4 (type C phenotype), and also found activity higher in the former rather than the latter. Thus, caution is needed when discussing prognosis in newly diagnosed patients with PCD regardless of enzymatic results. It means that PC activity measurement in skin fibroblast can prove if an unknown variant impairs PC activity, but not to which extent. Of note, all of our patients had germline mutations, and thus tissue mosaicism is unlikely to be playing a role in defining phenotype as has previously been suggested (Wang et al., 2008).

In the last years, in-silico free energy calculations have been increasingly used to estimate the effects of single amino acid mutations on protein stability and protein-ligand binding affinity (Gapsys et al., 2016; Fowler et al., 2018; Hauser et al., 2018). According to the results of our structural analysis, the close proximity of the residues G150, R451, A610, M743 (all associated to type A PCD) to the functionally relevant ligands may indicate that the respective variants could impair the interaction of the protein with the ligands (ATP, pyruvate, biotin and acetyl-CoA) or they could potentially alter the energetics of the catalytic carboxyl transfer steps. An example supporting the hypothesis that type A variants are linked with the disturbed interactions between the enzyme and its ligands comes from patient 2 described by Monnot et colleagues (Monnot et al., 2009): in this case, type A phenotype is associated with the p.R631Q and p.R270W variants.
p.R270W may interfere with the bicarbonate binding and biotin delivery to the BC domain.

According to our in-silico models, variants associated with the type C PCD do not directly interact with ATP, biotin and pyruvate. Some of these variants (p.S266A, p.S266Y, p.T569A) are located in the core of the BC and CT domains and destabilize the enzyme. S266 is neighboring C265 which has been speculated to be important for the CO₂ fixation from the bicarbonate (Li et al., 1992). This observation matches a positive response of the patient with the p.S266A variant to bicarbonate treatment (Van Coster et al., 1991). p.N1081S may have an effect on the interaction between the four PC monomers with each other or might affect the enzyme’s activation by acetyl-CoA.

For most of the described variants associated to both type A and type C PCD, free energy calculations predicted a destabilizing effect of the corresponding mutant amino acid on the monomer. Interestingly, for most of the missense variants associated to type A disease the balance between the two major conformational states (conformers 1 and 2) of the enzyme is disturbed; on the contrary, for all type C missense variants the balance between the two conformers appears to be unperturbed. Taken together, we hypothesize that a missense variant, which does not cause a precocious mRNA decay, leads to a mutant monomer with a residual enzymatic function and to the consequent mild type C phenotype if this mutant amino acid does not disturb the balance between the two major enzyme conformers. On the other hand, if the mutant amino acid leads to a misbalance between the two main conformers, the global enzymatic function of the mutant homotetramer is impaired to cause a severe type A phenotype.
5. CONCLUSION

This study improves our understanding of genotype-phenotype correlation in PCD. Considering the genetic data of all reported patients with type A and C phenotypes, we could observe that a late-onset mild PCD (type C) is not associated to homozygous variants but to heterozygous ones in PC. In general, our in-silico calculations of the PC free energy changes show that mutations which do not destabilize the balance between the two major enzyme conformers are likely to lead to the type C phenotype, while those mutations which affect balance between the two major conformers (regardless of predicted residual enzymatic activity) lead to a type A phenotype. Identification and characterization of additional PCD patients will help to further refine these predictions.

REFERENCES


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Schrödinger. The PyMOL Molecular Graphics System, Version 1.8.0.3, LLC.


FIGURES

FIGURE 1 Brain MRI. Coronal cut (A) and spectrographic peaks (B) from patient 2 (type A), two coronal cuts (C and D) from patient 3 (type A), one coronal cut from patient 4 (E) (type C). The arrows indicate the structural brain abnormalities.
FIGURE 2 Enzymatic activity of pyruvate carboxylase in skin fibroblasts. Enzymatic activity of pyruvate carboxylase was measured in skin fibroblasts of patients 2 and 3 (both type A) and patient 4 (type C) and was expressed as percentage value of the corresponding healthy control sample. One healthy control sample was used for each patient fibroblasts’ measurement.
FIGURE 3 Two states of the human pyruvate carboxylase. Conformer 1 (left) has its BCCP domain in the vicinity of BC domain, while in the conformer 2 (right) BCCP domain brings biotin to the CT domain. Both structures contain four homomonomers (chains A, B, C, D), of which monomer A is depicted in the cartoon representation, while three other monomers are in surface representation. The four domains of the monomer A are color coded: BC in yellow, CT in green, PT in violet, BCCP in pink. The ligands (ATP, biotin, CoA, pyruvate) are depicted as gray spheres. The novel mutated residues revealed in our study are shown in big colored spheres: for type A PCD the residues are depicted in red, for type C PCD in blue. The variants previously reported in literature are depicted in small colored spheres, either red for type A PCD or blue for type C PCD.

TABLE 1 Molecular characterization PC gene and protein in type A and type C PCD patients. Domains: biotin carboxylase (BC), carboxyl transferase (CT), PC tetramerization (PT), biotin carboxyl carrier protein (BCCP)

<table>
<thead>
<tr>
<th>Source</th>
<th>Patient</th>
<th>Amino acid variants</th>
<th>Involved protein domain(s)</th>
<th>Variant zygoty</th>
<th>Comments on mutant RNA's and protein's enzymatic activity</th>
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<td>this</td>
<td>1</td>
<td>p.G150</td>
<td>BC</td>
<td>homoz</td>
<td>ND</td>
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<table>
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<th>Control Sample</th>
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<td>CT</td>
<td>homozygous</td>
<td>reduced protein's import into mitochondria and faster degradation 2.3% of control sample</td>
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<tr>
<td>This study 3</td>
<td>p.R626 + unknown intronic variant</td>
<td>CT</td>
<td>compound heterozygous</td>
<td>allele 2: intronic variant causing disturbed pre-mRNA splicing 6% of control sample</td>
</tr>
<tr>
<td>Wexler et al, 1998</td>
<td>p.V145 A</td>
<td>BC</td>
<td>homozygous</td>
<td>barely detectable amount of protein 7-25% of control sample</td>
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<td>Monnot et al, 2009</td>
<td>p.R156 Q</td>
<td>BC</td>
<td>homozygous</td>
<td>ND</td>
</tr>
<tr>
<td>Wexler et al, 1998; Xiang et al, 2008</td>
<td>p.R451 C</td>
<td>BC</td>
<td>homozygous</td>
<td>disturbed activation by CoA 7% of control sample</td>
</tr>
<tr>
<td>Carbon et al, 2003;</td>
<td>p.A610 T</td>
<td>CT</td>
<td>homozygous</td>
<td>reduced protein's import into mitochondria and faster degradation 1-4% of control sample</td>
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<td>this study 4</td>
<td>p.S266 Y + unknown intronic variant</td>
<td>BC</td>
<td>compound heterozygous</td>
<td>allele 2: unknown intronic variant supposed to disturb pre-mRNA splicing</td>
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<tr>
<td>1-4 % of control sample</td>
<td></td>
<td></td>
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<td></td>
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<tr>
<td>p.M743 I</td>
<td>CT</td>
<td>homozygous</td>
<td></td>
<td></td>
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<tr>
<td>p.R270 W + p.R631 Q</td>
<td>BC and CT</td>
<td>compound heterozygous</td>
<td></td>
<td></td>
</tr>
<tr>
<td>p.R62C + p.A847 V + p.R631 Q</td>
<td>BC and CT</td>
<td>compound heterozygous</td>
<td></td>
<td>three allelic variants in a somatic mosaic constellation; mutant mRNA transcripts containing r.1892g&gt;a and r.2540c&gt;u were unstable and rapidly degraded</td>
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Type C Pa...
TABLE 2 Summary of free energy calculations for type A and type C variants. ΔΔG values denote changes in the PC stability calculated by thermodynamic modeling of each variant. ΔΔG conformer 1 and ΔΔG conformer 2 correspond to the changes in stability for the two modeled conformers. ΔΔΔG is the difference between the stability changes between the two conformers for each single variant, and represents the change in balance between the two major conformers in comparison to the WT enzyme. The free energy differences are reported as a range between the values estimated with two energy functions.

<table>
<thead>
<tr>
<th>Source</th>
<th>Variants</th>
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<th>ΔΔG\text{conformer1}, kcal/mol</th>
<th>ΔΔG\text{conformer2}, kcal/mol</th>
<th>ΔΔΔG, kcal/mol</th>
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<th>Val 2</th>
<th>Val 3</th>
<th>Val 4</th>
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<td>p.A610T</td>
<td>Homozygous</td>
<td>2.3 – 6.7</td>
<td>3.9 – 7.3</td>
<td>-1.6 – -0.6</td>
<td>5.5 – 8.0</td>
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<tr>
<td>This study</td>
<td>p.R626Q</td>
<td>Heterozygous</td>
<td>-12.4 – 3.4</td>
<td>-5.3 – 1.7</td>
<td>-7.1 – 1.7</td>
<td>0.0 – 1.8</td>
</tr>
<tr>
<td>Pineda et al, 1995; Wang et al, 2008</td>
<td>p.R62C</td>
<td>Heterozygous</td>
<td>-5.3 – 2.4</td>
<td>-4.2 – 0.7</td>
<td>-1.1 – 1.7</td>
<td>-3.1 – -1.0</td>
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<tr>
<td>Monnot et al, 2009</td>
<td>p.R156Q</td>
<td>Homozygous</td>
<td>2.5 – 50.2</td>
<td>-0.8 – 22.0</td>
<td>3.2 – 28.1</td>
<td>-6.1 – -4.0</td>
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<tr>
<td>Monnot et al, 2009</td>
<td>p.R270W</td>
<td>Heterozygous</td>
<td>-3.4 – 3.3</td>
<td>1.6</td>
<td>-5.0 – 1.7</td>
<td>0.0 – 6.6</td>
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<td>Wexler et al, 1998</td>
<td>p.R451C</td>
<td>Homozygous</td>
<td>-5.0 – 24.7</td>
<td>-2.0 – 11.4</td>
<td>-3.1 – 13.3</td>
<td>-1.9 – 1.1</td>
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<td>Wang et al, 2008</td>
<td>p.R631Q</td>
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<td>2.5 – 3.4</td>
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<td>p.M743I</td>
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<td>11.0 – 29.6</td>
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<td>7.3 – 14.7</td>
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<td>Wang et al, 2008</td>
<td>p.A847V</td>
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<td>1.4 – 2.2</td>
<td>1.4 – 1.7</td>
<td>-0.1 – 0.5</td>
<td>1.2 – 1.5</td>
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**Type C**

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<th>Study</th>
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<td>Wang et al, 1998; Van Coster et al, 1991</td>
<td>p.S266A</td>
<td>Heterozygous</td>
<td>-7.8 – 2.4</td>
<td>-5.0 – 1.3</td>
<td>-2.8 – 1.1</td>
<td>-2.2 – 0.2</td>
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<tr>
<td>Wang et al, 2008</td>
<td>p.T569A</td>
<td>Heterozygous</td>
<td>3.3 – 3.4</td>
<td>1.7</td>
<td>1.7</td>
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