Investigating the structural impacts of a novel missense variant identified with whole exome sequencing in an Egyptian patient with propionic acidemia

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\textbf{ABSTRACT}

Propionic Acidemia (PA) is an inborn error of metabolism caused by variants in the \textit{PCCA} or \textit{PCCB} genes, leading to mitochondrial accumulation of propionyl-CoA and its by-products. Here, we report a 2 year-old Egyptian boy with PA who was born to consanguineous parents. Biochemical analysis was performed using tandem mass spectrometry (MS/MS) on the patient's dried blood spots (DBS) followed by urine examination of amino acids using gas chromatography/mass spectrometry (GC/MS). Molecular genetic analysis was carried out using whole-exome sequencing (WES). The \textit{PCCA} gene sequencing revealed a novel homozygous missense variant affecting the locus (chr13:100962160) of exon 16 of the \textit{PCCA} gene, resulting in the substitution of the amino acid arginine with proline at site 476 (p.Arg476Pro). Computational analysis revealed that the novel variant might be pathogenic and attributed to decrease the stability and also has an effect on the biotin carboxylase c-terminal domain of the propionyl carboxylase enzyme. The physicochemical properties analysis using NCBI amino acid explorer study revealed restrictions in the side chain and loss of hydrogen bonds due to the variant. On the structural level, the loss of beta-sheet was observed due to the variant proline, which has further led to the loss of surrounding interactions. This loss of beta-sheet and the surrounding interactions might serve the purpose of the structural stability changes. The current study demonstrates that a combination of whole-exome sequencing (WES) and computational analysis are potent tools for validation of diagnosis and classification of disease-causing variants.

1. Introduction

Propionic Acidemia (PA) (MIM # 606054) is an autosomal recessive metabolic disorder caused by a deficiency in the activity of the propionyl-CoA carboxylase (PCC) enzyme. The PCC enzyme catalyzes the carboxylation of propionyl-CoA to methylmalonyl-CoA. It is encoded by the \textit{PCCA} (MIM# 232000) and \textit{PCCB} (MIM# 232050) genes to form a 750 kDa heterododecamer composed of α- and β-subunits, respectively [1]. Bi-allelic variants that diminish or abolish the function of \textit{PCCA} or \textit{PCCB} subunits result in accumulation of propionic acid and propionyl-CoA related metabolites, which are known to be toxic for the brain, heart, muscles, and liver [2,3]. Such accumulation of intermediate metabolites impacts the anaplerotic replenishment of TCA intermediates, the oxidative phosphorylation, and many other metabolic pathways [1].

Symptoms of PA may begin as early as the first day postpartum but can be delayed for months or years. For neonates, the classical clinical presentation consists of vomiting, dehydration, weight loss, temperature instability, neurological involvement with muscular hypo- or hypertonia, irritability, lethargy progressing to coma, seizures, and death, if untreated. However, symptoms after the neonatal period are non-specific and might mimic many other common disorders [4]. At presentation, laboratory findings include severe and persistent metabolic acidosis with an elevated anion gap, hyperammonemia, hyperglycinemia, and hyperglycinuria [4]. Diagnosis is made by mass spectrometry for acylcarnitines in the blood showed elevated C3-carnitine and C3-/C2-carnitine ratio, and gas chromatography for organic acids in urine showed increased in urine excretion of methyl citrate and 3-
hydroxy-propionate.

According to the Human Gene Mutation Database [5], a total of 207 variants in both PCCA and PCCB genes have been identified to date. The mutational spectrum is clearly population-specific, and some variants have been found to be exclusively associated with specific geographical or ethnic groups. For instance, the 1218del14ins12 (ins/del) is the most common variant among Caucasians [6,7]. The mutation p.T428I is reported frequently among Japanese and Korean PA patients [8–10]. For both PCCA and PCCB genes, missense variants are known to be the most frequent defects (43% and 58%, for PCCA and PCCB, respectively), followed by small insertions/deletions and splicing variants, most of them result in a truncated protein [5]. The variants are distributed all along with the gene sequences. However, some clustering is observed in exon 13 in the PCCA gene, and in exons 12 and 15 in the PCCB gene, implying that they could be hot spots for variants or involved in critical functional domains.

In this study we used NGS to identify a novel variant in a 2-year-old Egyptian boy with PA, who was born to consanguineous parents. This novel variant was further subjected to computational analysis to predict the pathogenicity and stability changes in the protein due to the occurrence of a novel variant in the PCCA gene.

2. Materials and methods

2.1. Ethics statement

This study was approved by the Institutional Review Boards (IRB) of Ain Shams University, and adhered to the ethical guidelines of the Declaration of Helsinki and its amendments. Informed consent was acquired from the parents of the proband after a detailed explanation of the purpose of the study.

2.2. Biochemical analysis

2.2.1. Tandem mass spectrometry (MS/MS)

A blood sample was taken from the patient through a finger prick on the Guthrie card (Whatman 903 filter paper (GE Healthcare, New Jersey, USA)). The blood spots were analyzed for acylcarnitines and amino acids by a triple-quadrupole tandem mass spectrometer (ACQUITY UPLC H-Class. Waters® corporation, Massachusetts, USA), with a positive electrospray ionization probe, utilizing Mass Chrom® Amino acids and Acylcarnitines from Dried Blood kit (Chromsystems Instruments & Chemicals GmbH, München, Germany) according to manufacturer's instructions. The data of the MRM scan were analyzed using Neolynx® application (Waters® Corporation, Massachusetts, USA).

2.2.2. Gas chromatography/ mass spectrometry (GC/MS)

A urine sample was collected from the patient and frozen until derivatization by the silylation of organic compounds. The GC/MS analysis was done using the Agilent system Gas chromatography instrument that is interfaced with a Mass spectrometer and Gas chromatography capillary column (Agilent, USA).

2.3. Molecular genetic analysis

2.3.1. Next-generation sequencing

Peripheral whole blood samples were collected from the patient and parents. DNA was extracted (using Gene JET Whole Blood DNA purification Mini Kit (Thermo Scientific, Germany). The initial amount of whole genomic DNA was amplified using the Ampliseq RDY panel kit (Thermo Fisher Scientific, Waltham, MA USA 02451). Read trimming, alignment (hg19), and variant calling were automatically completed using the Ion Torrent Version 5.6 (Thermo Fisher Scientific, Waltham, MA USA). Variant calling files (.vcf) were uploaded to the Ion Reporter System (Thermo-Fisher Scientific, Waltham, MA, USA) for variant annotation. Sequence alignment and extraction of SNPs were performed.

2.3.2. Sanger sequencing

Sanger sequencing was performed to identify variants identified by WES and to determine the co-segregation of the candidate variant with the disease in the parents. PCR amplification was performed, and primer pairs were designed using Primer3 software (http://primer3.ut.ee/). Sequencing reactions of PCR products were carried out using BigDye Terminator v3.1 chemistry and separated on ABI 3500 genetic analyzer (Applied Biosystems) according to manufacturer's instructions. GenBank RefSeq NM_000282.2 was used as the PCCA reference sequence.

2.4. Computational analysis

2.4.1. Sequence and structural dataset

The protein sequence for the PCCA was obtained from the UniProt database with the ID P05165 [11]. The PCCA protein lacks the crystal structure in the public databases to date. Henceforth, the protein sequence obtained from the UniProt database was used to build the 3D structure of the protein using the Swiss Model online server [12]. The 3D protein model was verified using the online RAMPAGE server [13] based on an assessment of the Ramachandran Plot [14]. The RAMPAGE server approves the protein structure on the premise of φ, ψ point of individual deposits [15,16].

2.4.2. Pathogenicity, stability, physicochemical properties analysis

The pathogenicity and the effect of the variant in destabilizing the protein structure were studied using a series of algorithms. PolyPhen-2 [16], fathmm [17], PhD-SNP [18], Pmut [19], and Pon-P2 [20], were used to predict the pathogenic effect of the novel variant p.Arg476Pro. The destabilizing properties of the p.Arg476Pro variant were studied using the mCSM [21], SDM [22], DUET [23], ENCoM [24], and DynaMut algorithms [25]. The change in physicochemical properties of the native and variant amino acids was studied using the NCBI amino acid explorer [26].

2.4.3. Variant modeling and structural analysis

The novel variant p.Arg476Pro was introduced in the modeled native structure using SwissPDB viewer, and the structure was energy minimized using the same [27,28]. Visualization of the proteins and computation of the global main chain root-mean-square distance (RMSD) between the native and mutant protein was performed using Discovery Studio Visualizer software v20.1.0.19295 (Dassault Systèmes BIOVIA, San Diego, CA, USA). The change in surrounding amino acids and hydrogen bonds was also studied using the same software.

3. Results

3.1. Patients characteristics

The proband was a 2-year-old Egyptian boy, the second child of first-cousin consanguineous marriage. He was born at the 40th week of gestation, by spontaneous vaginal delivery, with a birth weight of 3.5 Kg. The family history record showed a previously deceased sibling.

The first child of the couple was a female born at full-term by spontaneous vaginal delivery, with a birth weight of 2.75 Kg. She had poor suckling and developed jaundice on day 3. She was admitted to NICU, where breast feeding was stopped, and she received phototherapy and protein-free parenteral feeding. After a few days, she was improved and discharged. At the age of 4 month, she was readmitted for lethargy, sleepiness, poor sucking, and vomiting after every feed. She was diagnosed with sepsis and died (Fig. 1).

The proband presented at the age of 3 days with jaundice and was admitted to NICU for management. Blood and urine samples were
infection and associated with marked hyperammonemia (412 μg/dL), hypoglycemia, and acidosis. Follow-up of biochemical studies at one year of treatment showed that the level of ammonia was 81 μg/dL, C3-carnitine was 17.71 μmol/L, and C3: C2 ratio was 0.79.

### 3.2. Biochemical findings

The laboratory analysis showed a high level of total bilirubin (14.8 mg/dL), mostly unconjugated bilirubin (11.9 mg/dL). The hyperbilirubinemia was accompanied by metabolic acidosis and an elevation of ammonia (358 μg/dL) and lactate (51.4 mg/dL) (Table 1). The biochemical analysis of the amino acid-acylcarnitine was performed on a dried blood spot (DBS) using LC-MS/MS technique, and findings were consistent with either propionic or methylmalonic acidemia. The MS/MS measurement revealed highly elevated propionyl carnitine (C3, 27.34 μmol/L) with a very highly elevated C3:C2 ratio (0.41) (Table 1).

Urine examination for organic acid using GC/MS showed a moderately elevated hydroxy propionic and significantly elevated methyl citrate that were suggestive of PA. The huge amount of hippuric acids was masking some other analytes.

### 3.3. Molecular genetic findings

Whole exome sequence analysis showed the patient to be homozygous for a novel missense variant affecting the locus (chr13:100962160) of exon 16 of the PCCA gene. The c.1427G > C mutation leads to the substitution of the amino acid Arginine with Proline at site 476 of the amino acid sequence of the protein chain (p.Arg476Pro). No other pathological or likely pathological variants were detected in the PCCA and PCCB loci. This variant was absent in 300 normal chromosomes. Sanger DNA sequencing confirmed the heterozygous status of both parents for the same variant.

### 3.4. Computational analysis

#### 3.4.1. Protein modeling and validation

The 3D structure of the PCCA protein was modeled using the online Swiss Model web-based client. The crystal structure of the holoenzyme of propionyl-CoA carboxylase (PCC) of the Ruegeria pomeroyi species with PDB ID 3N6R was found to be the most suitable template to build the protein model with a sequence identity of 53.94%. This modeled protein structure was further subjected to the validation for Ramachandran plot calculations. This validation was done using the online RAMPAGE server. From the validation plot, we observed that the number of residues in the favored region was 89.0%, the number of residues in the allowed region was 8.4%, and the number of residues in the outlier region was 2.6% (Fig. 2).

#### 3.4.2. Pathogenicity, stability, physicochemical properties analysis

Pathogenicity and stability, physicochemical properties analysis

Pathogenicity and stability of the novel variant p.Arg476Pro were predicted using five algorithms for each. The pathogenicity predictors, PolyPhen-2, FATHMM, PhD-SNP, Pmut, and Pnom–P2, predicted the new variant p.Arg476Pro to be probably damaging, damaging, disease, and pathogenic, respectively (Table 2). The stability predictors, mCSM, SDM, DUET, ENCoM, and DynaMut predicted the new variant p.Arg476Pro to be destabilizing (Table 2). The difference in the physicochemical properties of the native arginine and variant proline was studied using the NCBI amino acid explorer (Table 3). From the study, it was identified that the side-chain flexibility, which was high in the native arginine, was restricted due to the variant proline. Similarly, due to the variant proline, the Ionic and H-bonds were lost, which were initially present in the native arginine. Significantly, the loss of 7 potential side-chain H-bonds due to the variant proline was also observed (Table 4).

**Table 1**

| Analytes                         | Result  | Reference Range |
|----------------------------------|---------|-----------------|
| Total bilirubin                  | 14.8 mg/dL | < 1 mg/dL      |
| Direct bilirubin                 | 2.9 mg/dL  | < 0.25 mg/dL     |
| Indirect bilirubin               | 11.9 mg/dL | < 0.75 mg/dL     |
| Lactate                          | 51.4 mg/dL | 4.5–20 mg/dL    |
| Ammonia                          | 358 μg/dL  | 11–102 μg/dL     |
| Acetylcarnitine (C2)             | 27.34 μmol/L | 5–47 μmol/L    |
| Propionylcarnitine (C3)          | 10.81 μmol/L | 0–7 μmol/L     |
| C3/C2 ratio                      | 0.41      | 0.25            |

Fig. 1. Family Pedigree, demonstrating that the parents of the proband were heterozygous for the p.Arg476Pro variant.
3.4.3. Structural analysis

The structural analysis of the native and PCCA with variant p.Arg476Pro proteins was studied using Discovery Studio Visualizer software v20.1.0.19295 (Dassault Systèmes BIOVIA, San Diego, CA, USA). The RMSD between the native and variant PCCA was found to be 0.038 Å. The loss of beta-sheet due to the variant proline was also observed (Fig. 3). In addition, we observed the loss of interacting amino

![Diagram](Image)

**Fig. 2.** Ramachandran plot validation of the modeled PCCA 3D protein structure. The plot reveals that the number of residues in the favored region was 89.0%, the number of residues in the allowed region was 8.4%, and the number of residues in the outlier region was 2.6%.

**Table 2**

| S/No | Algorithm | Prediction      | Score  |
|------|-----------|-----------------|--------|
| 1    | PolyPhen-2| Probably damaging| 1      |
| 2    | fathmm    | Damaging        | −1.55  |
| 3    | PhD-SNP   | Disease         | 1      |
| 4    | Pmut      | Disease         | 0.71   |
| 5    | Pon-P2    | Pathogenic      | 0.9    |

**Table 3**

| S/No | Algorithm | Prediction  | ΔΔG (kcal/mol) |
|------|-----------|-------------|----------------|
| 1    | mCSM      | Destabilizing| −0.109         |
| 2    | SDM       | Destabilizing| −1.54          |
| 3    | DUET      | Destabilizing| −0.489         |
| 4    | ENCoM     | Destabilizing| −0.595         |
| 5    | DynaMut   | Destabilizing| −0.01          |

**Table 4**

The difference in the physicochemical properties of native arginine and variant proline of the novel variant p.Arg476Pro of PCCA protein.

| Feature                  | Arginine | Proline |
|--------------------------|----------|---------|
| Side-chain flexibility   | High     | Restricted |
| Interaction modes        | Ionic, H-bonds, van der Waals | van der Waals |
| Potential side-chain H-bonds | 7 | 0 |
| Residue molecular weight | 156      | 97      |
| Isoelectric point         | 10.8     | 6.3     |
| Hydrophobicity            | 0        | 0.678   |
| Standard codon(s)         | CGN, AGR | CCN     |
| Charge                    | Positive | Nonpolar |

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acids and hydrogen bonds due to the novel variant p.Arg476Pro. The native arginine at position 476 showed four amino acids interactions (SER418, GLN419, ILE475, and GLN615) with six conventional hydrogen bonds (2 with GLN415, 2 with GLN615, 1 with ILE475, and 1 with SER418) and one covalent bond (ILE475) (Fig. 4A). While the variant proline at position 476 showed only two amino acid interactions (SER418 and GLN419) with one conventional hydrogen bond, respectively (Fig. 4B).

4. Discussion

In this study, we report the identification of a novel homozygous variant PCCA: c.1427G > C (p.Arg476Pro), affecting the locus (chr13:100962160) of exon 16 of PCCA gene, in an Egyptian patient diagnosed with propionic acidemia; the diagnosis was initially made according to the clinical evaluation, acylcarnitine, and organic acid profiles and confirmed by next-generation sequencing of the patient and Sanger sequencing of his parents who were found to be heterozygous for the same variant. To the best of our knowledge, this variant has not been reported in the current human genome databases (1000 Genomes, dbSNP, ESP, ExAC, HGMD, and gnomAD), and was absent in 300 normal chromosomes.

History of consanguinity and the neonatal death of the sister of the proband led us to do a careful genetic investigation, especially with the classical presentation of propionic acidemia for his deceased sister. Although she was diagnosed with neonatal sepsis, the diagnosis of propionic acidemia should also be considered based on her clinical presentation and her transient improvement upon discontinuation of oral feeding during hospitalization. Advancement of high-throughput sequencing, particularly WES, has facilitated identifying variations or even novel genes associated with human diseases and complex traits. In this study, used WES to identify this novel variant PCCA: c.1427G > C, which was found in the proband with a WES sequencing coverage depth of 100×, which was convinced to call the variant. No variants were found in the PCCB gene. The heterozygous status was confirmed in both parents using Sanger sequencing.

In the absence of newborn screening, the diagnosis of several inborn errors of metabolism at primary care units is likely to be missed or delayed; indicating the importance of the availability of a nationwide extended screening program to prevent the common morbidities and mortalities of patients with an inborn error of metabolism especially in developing countries [7]. The clinical application of our study is the confirmation of the genetic diagnosis in patients, enabling accurate genetic counseling and, in many cases, providing a prognostic view of the probable course of the disease. However, the genetic heterogeneity present in PA patients, with most variants being "private," present in only one family, along with the practical absence of homozygosity among the studied patients, hinders a straightforward genotype-phenotype correlation [7,29], and, in other cases, modulation of the effect of particular variants by genetic and epigenetic factors contribute to phenotypic variability.

The associated metabolic acidosis resulting from the accumulation of branched-chain amino acid, their intermediates, and all upstream metabolites of propionyl-CoA and hyperlactatemia is related to the potential indirect influence of propionyl-CoA to decrease the activity of pyruvate dehydrogenase and increase lactate [6]. In the presence of hyperammonemia, determination of blood amino acids, acylcarnitine, and the urinary organic acid using LC-MS/MS and GC/MS techniques were essential to lead for accurate differential diagnosis toward PA.

The PCCA with p.Arg476Pro variant was found to be pathogenic by all the algorithms utilized in this study (Table 2). According to the standards and guidelines for the interpretation of sequence variants and the joint consensus of the American College of Medical Genetics and the Association of Molecular genetics, this variant is "likely pathogenic (V)"
It fulfills 2 moderate and 3 supporting criteria, namely PM1, PM2, PP2, PP3, and PP4.

The stability predictors suggest that the variant-induced in protein might lead to destabilization of the PCCA protein structure (Table 3). This is a common mechanism in many metabolic diseases [30,31]. They are also known to play a crucial role in maintaining the stability of the protein [32–36]. Proline amino acid is well studied as a potent secondary structure breaker of alpha-helices as well as the beta sheets [37]. In the structural study, we observed a transformation of beta-sheet to loop, thus serving a main reason for the loss of stability, which correlates with our stability predictors’ results (Fig. 2 and Table 3). Furthermore, amino acid interactions and hydrogen bonds are known to play a crucial role in maintaining the stability of the proteins [38,39]. In our study, the loss of hydrogen bonds and the interacting amino acids revealed by the structural analysis further elucidate that the protein could have severe stability loss due to the novel variant p.Arg476Pro. This stability loss and protein structural changes make this novel variant more significant in causing PA. These findings agree with previous studies indicating that the loss of function of most PCCA variant proteins is based on protein reduction due to decreased stability [40,41].

The distinctive architecture among Arabs and the common consanguineous marriage practices is expected to create a unique susceptibility profile for rare genetic diseases, including PA [32]. Sanguineous marriage practices is expected to create a unique susceptibility profile for rare genetic diseases, including PA [32].

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