Research Article

Propionic and Methylmalonic Acidemias: Initial Clinical and Biochemical Presentation

Amira Mobarak,1 Heba Dawoud,1 Wesam A. Mokhtar,2 Abdelrahim A. Sadek,3 Gihan Mohamed Bebars,4 Amr Ahmed Othman,5 Rofaida M. Magdy,5 Hanaa Nofal,6 and Amr Zoair7

1Medical Biochemical Diseases Division, Pediatrics Department, Faculty of Medicine, Tanta University, Egypt
2Pediatrics Department, Faculty of Medicine, Zagazig University, Egypt
3Neurology Division, Pediatrics Department, Faculty of Medicine, Sohag University, Egypt
4Pediatrics Department, Minia University, Egypt
5Medical Genetics Division, Pediatrics Department, Faculty of Medicine, Sohag University, Egypt
6Clinical Pathology Department, Faculty of Medicine, Tanta University, Egypt
7Pediatrics Department, Faculty of Medicine, Tanta University, Egypt

Correspondence should be addressed to Amira Mobarak; dr.amiramobarak@gmail.com

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PA and MAA have numerous nonspecific presentations, potentially leading to delayed diagnosis or misdiagnosis. In this paper, we present the clinical and biochemical characteristics of MMA and PA patients at initial presentation. Results. This is a retrospective review of 20 patients with PA (n = 10) and MMA (n = 10). The most observed symptoms were vomiting (85%) and refusing feeding (70%). Ammonia was 108.75 ± 9.3 μmol/l, showing a negative correlation with pH and bicarbonate and positive correlation with lactate and anion gap. Peak ammonia did not correlate with age of onset (r = 0.11 and p = 0.64) or at diagnosis (r = 0.39 and p = 0.089), nor did pH (r = 0.01, p = 0.96; r = −0.25, p = 0.28) or bicarbonate (r = 0.07, p = 0.76; r = −0.22, p = 0.34). There was no correlation between ammonia and C3 (r = 0.1 and p = 0.96) or C5 (r = 0.23 and p = 0.32). The glycine was 386 ± 167.1 μmol/l, and it was higher in PA (p = 0.003). There was a positive correlation between glycine and both pH (r = 0.56 and p = 0.01) and HCO3 (r = 0.49 and p = 0.026). There was no correlation between glycine and ammonia (r = −0.435 and p = 0.055) or lactate (r = 0.32 and p = 0.160). Conclusion. Clinical presentation of PA and MMA is nonspecific, though vomiting and refusing feeding are potential markers of decompensation. Blood gas, lactate, and ammonia levels are also good predictors of decompensation, though increasing levels of glycine may not indicate metabolic instability.

1. Introduction

Propionic acidemia (PA) and methylmalonic acidemia (MMA) are autosomal recessive diseases. They are caused by a defect in propionyl-CoA carboxylase and methylmalonyl-CoA enzymes, respectively.

They present classically in an “early-onset” neonatal acute life-threatening form, including lethargy, vomiting, shock, dehydration, metabolic acidosis, and hyperammonemia. This acute presentation is known as “acute decompensation” which can be fatal if not treated properly. PA and MMA can also present later in life with chronic and more insidious onset. While more common in early onset, both suffer from acute decompensation episodes throughout the course of the illness. These episodes are usually triggered by infection, fasting, fever, or any other stress condition [1, 2].

A wide array of presenting symptoms, which are almost always nonspecific, can lead to delayed diagnosis or misdiagnosis. Multiple reports have addressed the primary presentation of patients with PA and MMA, each describing different symptoms overall as well as different most frequent symptoms [3–11], including atypical neurological presentation [12–14].
This paper reports the clinical and biochemical characteristics of 20 Egyptian patients with MMA and PA at first presentation, with the hopes of enhancing knowledge surrounding the presenting signs and symptoms of these two conditions.

2. Patients and Methods

We reviewed 20 Egyptian patients with PA (n = 10) and MMA (n = 10) diagnosed from year 2014 to 2018. This study was approved by the ethics committee of the Faculty of Medicine, Tanta University, Egypt. Written informed consent was obtained from the patients’ legal guardians/parents. All data was collected retrospectively from patient charts.

2.1. Statistical Analysis. The data collected and inserted into a Microsoft Excel spreadsheet for analysis are presented in the form of mean ± standard deviation (SD). The Shapiro-Wilk test was used to check for data normality, Mann-Whitney, Student’s t-test, and Fisher’s exact tests were used to compare data groups. Pearson’s linear correlation was used to detect the relationship between different variables. A multiple regression analysis was calculated to predict the ammonia level of the cohort based on the lactate and bicarbonate levels. Significance value was set at p < 0.05.

3. Results

3.1. Clinical Presentation. This study included 20 patients, PA (n = 10) and MMA (n = 10) with age 3.99 ± 1.29 years at the time of data collection. Patients were diagnosed by selective screening prompted by positive family history (one patient) or suspicious preliminary investigations and clinical presentation. All patients were of a Middle Eastern background and born at term (m = 38.6 ± 2.18); 80% had a history of consanguinity, and 45% were females. The majority of patients (70%) were delivered by spontaneous vaginal delivery. The birth weight was 3.9 ± 1.2 kg, with no significant difference between PA (3.5 ± 1.33 kg) and MMA (4.4 ± 1.17 kg) (p = 0.16; two tail).

The age of onset of symptoms was 5.38 ± 1.38 months, with 60% of patients presenting in the first 3 months of life, only three (15%) in the neonatal period, and 40% after 3 months of age with no significant difference between PA (4.35 ± 6.29 months) and MMA (6.32 ± 6.25 months), and p = 0.49. The age at diagnosis was 8.05 ± 7.6 months, with no significant difference between MMA (8.7 ± 6.8 months) and PA (8.05 ± 8.8 months) (p = 0.85; two tail). The time to diagnosis after the onset of initial symptoms was 3.03 ± 5.5 months, with no difference between PA (3.7 ± 7.6 months) and MMA (2.37 ± 2.26 months) (p = 0.6; two tail), or between early onset (3.8 ± 6.8 months) and late onset (1.77 ± 2.5 months) (p = 0.14; two tail).

Almost all patients, regardless of age at diagnosis, presented with acute metabolic decompensation apart from one patient who presented with acute ataxia without signs of metabolic derangement. The most frequent presenting symptoms were vomiting (85%) followed by refusing feeding (70%), dehydration (60%), encephalopathy (55%), hypotonia (55%), shock (45%), hepatomegaly (45%), seizures (20%), dystonia (10%), and ataxia (5%). There was no significant statistical difference between PA and MMA in frequency of presenting symptoms, (p > 0.05) except for hypotonia which was more frequent in PA (p = 0.038; two tail). The triggering factors were fever and GI illness in 45% of the patients and upper respiratory infection in 40%, and no trigger factor could be identified in 15%.

3.2. Biochemical Presentation. The pH was 7.32 ± 0.06. The standard bicarbonate was 12.2 ± 5.71 mmol/l, with no significant difference between PA (12.9 ± 5.67 mmol/l) and MMA (11.4 ± 5.76 mmol/l) (p = 0.038). The anion gap was 22.5 ± 8.38, and the MMA patients tended to have higher anion gap, with p = 0.005 (Tables 1 and 2). Nineteen patients (19/20) had positive ketones in their urine with a median of +2, with no significant difference between the PA and MMA groups (p = 0.53).

Ammonia level, however, had significant negative correlations with both pH and bicarbonate levels and a positive correlation with lactate and anion gap (Figure 1). A multiple regression analysis was calculated to predict the ammonia level of the cohort based on the lactate and bicarbonate levels. The results of the regression suggest that level of lactate and bicarbonate can predict 67% of the variance (r² = 0.67, F (2.17) = 16.99, and p < 0.001). It was found that lactate significantly predicted ammonia level (β = 0.52 and p = 0.004), as did bicarbonate (β = −0.47 and p = 0.008), at a confidence interval of 95%.

Lactate level was 2.87 ± 1.53 mmol/l, where (11/20) patients had lactate above 2.2 mmol/l. Interestingly, lactate did not show significant correlation with alanine (r = −0.3 and p = 0.14).

Glycine level was 386 ± 167.1 μmol/l and tended to be significantly higher in PA (p = 0.003). There was a positive correlation between glycine and both pH (r = 0.56 and p = 0.01) and HCO₃ (r = 0.49 and p = 0.026). No significant correlation was found between glycine and both ammonia (r = −0.435 and p = 0.055) and lactate (r = 0.32 and p = 0.160) (Figure 2).

MMA patient age at diagnosis was 8.7 ± 6.8 months, with all patients presenting after one month of life. The MMA group had a C3 : C2 of 0.75 ± 0.52 and a C3 of 14.3 ± 5.36 μmol/l. PA patient age at diagnosis was 8.05 ± 2.7 months, with a C3 value of 17.6 ± 3.99 μmol/l and C3 : C2 of 1.11 ± 0.14. Although C3 was higher in the PA group, it was not statistically significant (p = 0.14), and the C3 : C2 ratio did not show any significant difference between both groups (p = 0.06; two tail).
Table 1: Laboratory parameters at initial presentation.

| Lab. parameter      | Range (SI)                  | Mean ± SD                  | Normal reference range |
|---------------------|-----------------------------|----------------------------|------------------------|
| pH                  | 7.23-7.43                   | 7.32 ± 0.06                | 7.35-7.45              |
| HCO₃⁻                | 4.25                        | 12.8 ± 5.71 mmol/l         | 22-26                  |
| Anion gap            | 10-39                       | 22.5 ± 8.38                | 12-15                  |
| Ammonia              | 45-212                      | 108.75 ± 41.82 μmol/l      | 9-33 μmol/l            |
| Lactate              | 1.1-6.5                     | 2.87 ± 1.53 mmol/l         | 1-2.2 mmol/l           |
| Blood glucose        | 67-250 (3.7-13.8 mmol/l)     | 156.65 ± 44.21 mg/dl (8.7 ± 2.47 mmol/l) | 70-126 mg/dl |
| Uric acid            | 2.5-11.2 (148.71-666.23 μmol/l) | 4.45 ± 1.85 mg/dl (264.7 ± 110.04 μmol/l) | 3.5-4.5 mg/dl |
| Glycine              | 77.5-721                    | 386.56 ± 167.11 μmol/l     | 20-500 μmol/l          |
| Alanine              | 99.56-401                   | 245.97 ± 88.65 μmol/l      | 20-600 μmol/l          |
| C3                   | 9.98-27.7                   | 16.06 ± 4.89 μmol/l        | 0-7 μmol/l             |
| C3 : C2              | 0.31-1.68                   | 0.93 ± 0.42 μmol/l         | 0-0.25 μmol/l          |
| Creatinine           | 0.4-0.8 (35.37-70.74 μmol/l) | 0.62 ± 0.12 mg/dl (54.82 ± 10.61 μmol/l) | 0.6-1 mg/dl |
| AST                  | 17-74                       | 41.45 ± 14.06 U/l          | 20-45                  |
| ALT                  | 20-69                       | 37.7 ± 13.41 U/l           | 20-45                  |
| Albumin              | 29-45                       | 35.8 ± 4.8 g/l             | 35-50 g/dl             |
| Hemoglobin           | 87-120                      | 107 ± 10.4 g/l             | 100 – 115 g/l          |
| Platelets            | 157,000-470,000             | 327,950 ± 101,710          | 150 – 450 × 10⁹/mm³    |
| White blood cells    | 2700-7000                   | 4490 ± 1121.98             | 3,500-5,000/mm³        |

Table 2: Comparison between the initial laboratory parameters of PA and MMA patients.

| Lab parameter      | PA Mean ± SD (SI) | MMA Mean ± SD (SI) | p value |
|--------------------|-------------------|--------------------|---------|
| pH                 | 7.34 ± 0.05       | 7.3 ± 0.06         | 0.14    |
| HCO₃⁻ (mmol/l)     | 14.84 ± 5.8       | 10.76 ± 4.97       | 0.11    |
| Anion gap          | 17.6 ± 6.13       | 27.42 ± 7.5        | 0.005*  |
| Ammonia (μmol/l)   | 107.4 ± 44.39     | 110.1 ± 41.44      | 0.70    |
| Lactate            | 2.52 ± 0.82       | 3.22 ± 1.99        | 0.70    |
| Blood glucose (mg/dl) | 154.7 ± 47.12 (8.5 ± 2.5 mmol/l) | 158.6 ± 43.55 (8.8 ± 2.4) | 0.84    |
| Uric acid (mg/dl)  | 4.01 ± 0.56 (238.5 ± 33.31 μmol/l) | 4.88 ± 2.55 (290.28 ± 151.68 μmol/l) | 0.81    |
| Glycine (μmol/l)   | 490.26 ± 158.8    | 282.84 ± 99.08     | 0.0032* |
| Alanine (μmol/l)   | 243.11 ± 84.42    | 248.83 ± 97.19     | 0.88    |
| C3 (μmol/l)        | 17.69 ± 3.94      | 14.43 ± 5.36       | 0.14    |
| C3 : C2            | 1.11 ± 0.14       | 0.75 ± 0.52        | 0.06    |
| Urea (mg/dl)       | 28.5 ± 11.21 (4.74 ± 1.86 mmol/l) | 23.3 ± 2.58 (3.87 ± 0.47) | 0.10    |
| Creatinine (mg/dl) | 0.65 ± 0.09 (57.46 ± 7.95 μmol/l) | 0.58 ± 0.13 (51.27 ± 11.49 μmol/l) | 0.36    |
| AST (U/l)          | 47.8 ± 16.41      | 47.8 ± 35.1        | 0.04*   |
| ALT (U/l)          | 40.2 ± 17.67      | 35.2 ± 7.28        | 0.42    |
| Albumin (g/l)      | 33.3 ± 4.8        | 38.3 ± 3.2         | 0.01*   |
| Hemoglobin (g/l)   | 114.3 ± 16.3      | 105.7 ± 10.9       | 0.22    |
| White blood cells (cells/mm³) | 4280 ± 1444.37 | 4700 ± 687.99 | 0.42 |
| Platelets × 10⁹/mm³| 330.4 ± 105.12    | 325.5 ± 103.79     | 0.91    |

*Significance value p < 0.05.
The mean level of uric acid was 4.45 ± 1.85 mg/dl (264.7 ± 110.04 μmol/l). One MMA patient had uric acid of 11.2 mg/dl (666.23 μmol/l) and was on allopurinol for four days. Mean level of random blood glucose (RBG) was 156.6 ± 44.2 mg/dl (8.7 ± 2.47 mmol/l); in 15/20 (75%), it was >126 mg/dl (7 mmol/l) (Table 3). Mean albumin level was 35.8 ± 4.7 g/l. It tended to be lower in PA (33.3 ± 4.8 g/l) when compared to MMA (38.3 ± 3.3 g/l and \( p = 0.014 \)). Five patients had a white blood cell count \( \leq 3,500 \) cells/mm\(^3\), and one patient had platelets of 157,000/mm\(^3\). Six patients had hemoglobin of <10.5 g/dl.

4. Discussion

The study is aimed at describing the clinical and biochemical presentations of MMA (\( n = 10 \)) and PA (\( n = 10 \)) in an Egyptian cohort of patients. All patients in this study were born at full term (GA > 37 weeks) or decrease in birth weight. This is in contrast with Kolker et al. [6] who noted that patients with MMA tended to have low birth weight, hence postulating that there could be a potential intrauterine disease process in the form of intrauterine impaired energy metabolism. Interestingly, PA patients in their cohort did not have low birth weight.

In our cohort, most patients (55%) presented in the first 3 months of life and only 15% presented in the neonatal period. This is in contrast to previous studies that demonstrated a higher frequency of patients presenting in the neonatal period [3, 5, 6, 10]. Sass et al. reported that 92% of their patients presented in the first 90 days of life, with 74% of their early-onset group presenting within the first 8 days of life and 26% between 11 and 90 days [9]. It was previously reported that patients with late onset tended to have a more obvious diagnostic time delay [6]. However, there was no correlation between the diagnostic time delay and age of onset in our cohort.

Previous studies show patients with late onset commonly presenting with chronic neurological symptoms, developmental delay, or organ damage rather than acute metabolic
decompensation (which was reported to be more common in early onset) [6, 10]. However, all patients in our cohort presented with metabolic decompensation of different severities regardless of age of onset, except one patient who presented with acute ataxia without any signs of metabolic decompensation. In addition, the neurological symptoms of seizures and dystonia were in the context of an episode of metabolic decompensation.

We must acknowledge that this is a retrospective study, and the age of onset of symptoms was concluded based on medical records and history taking. It is possible that earlier, more subtle symptoms, which may have in fact been more accurate indicators of age of onset, were not recorded through these means; obviously only major events, such as admissions and ER visits, can be reliably captured, while nonmajor events, which may still cause significant clinical burden, are more difficult to ascertain. In addition, genotype is one of the factors that can help establish severity, a factor that is unfortunately unavailable for our cohort.

The most commonly observed initial symptoms in our cohort were vomiting and refusing feeding, indicating that these are major clinical signs [15, 16] and should alert the physician of possible decompensation. We did not note any significant difference between PA and MMA patients with regard to the frequency of presenting symptoms, with the exception of hypotonia which tended to be more frequent in PA. Hypotonia was described previously as a frequent neurological complication of PA whether at initial presentation or later on [2].

We noted that ammonia had a negative correlation with pH and bicarbonate levels and a positive correlation with the anion gap and lactate. These biochemical abnormalities reflect secondary metabolic changes induced by accumulating toxic metabolites/organic acids including propionic acid. These metabolites inhibit PDH and TCA cycle enzymes and decrease succinyl-CoA production. In addition, they inhibit CPS I enzyme either directly or indirectly through N-acetyl glutamate synthase inhibition [17, 18]. We also observed that bicarbonate and lactate can be good predictors of hyperammonemia. By using these three markers together works to determine a more specific decompensation state of the patient [15, 16].

Unlike [6] who noted that late-onset patients had more severe metabolic acidosis compared to early-onset patients, with early-onset patients having higher ammonia levels, in our cohort, ammonia and pH did not correlate with age of onset or age at diagnosis. Interestingly, we found a positive correlation between glycine level and both pH and bicarbonate level, aligning with the hypothesis postulated by Al-Hassnan et al. [19]. They postulated that the conjugation of propionate with glycine or carnitine for excretion is pH sensitive, and, at normal acid-base status, propionyl carnitine might be the preferred mechanism; this is in contrast with metabolic acidosis when propionyl glycine might be the predominant form of propionate excretion.

![Figure 2: Correlation between the glycine in μmol/l on x-axis and (a) pH and (b) the standard bicarbonate levels.](image-url)
5. Conclusion

Clinical presentation of PA and MMA can vary, presenting a hurdle for timely diagnosis and management. For any patient who presents with nonspecific symptoms in the context of a minor metabolic stress, a metabolic disorder should be suspected. Vomiting and refusing feeding are also “red flags” to potential decompensation. In addition, our data indicates that blood gas, lactate, and ammonia levels are good predictors of acute decompensation when used in combination with patients’ biochemical evaluation. Increasing levels of glycine might not be an indicator of metabolic instability in such patients.

Data Availability

The datasets generated and/or analyzed during the current study are available from the corresponding author upon reasonable request.

Additional Points

Take Home Message. PA and MMA have nonspecific presenting symptoms, though vomiting and refusing feeding are “red flags” for decompensation. Moreover, blood gas and lactate taken with ammonia level are also good markers for biochemical instability.

Ethical Approval

This study has been approved by the ethics committee of the Faculty of Medicine, Tanta University, Egypt.

Consent

A signed informed consent was collected from the parents/-caregivers of all the participants in this study.

Conflicts of Interest

All the authors of this paper have no conflict of interest and nothing to declare.

Authors’ Contributions

Amira Mobarak conceived the presented idea, collected the data, analyzed the data, and wrote the manuscript. Heba Dawoud reviewed the results and the manuscript. Wesam A. Mokhtar, Abdelrahim A Sadek, Amr Ahmed Othman, Gihan Bebars, Rofaida M. Magdy, and Hanaa Nofal contributed to the data collection and manuscript revision. Amr Zoair supervised the conception of the idea and reviewed the final data analysis and manuscript.

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