The Effect of Carnitine Supplementation on Blood Ammonia Level in Epilepsy Children Treated with Valproic Acid: A Randomized Controlled Trial

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Background: Long-term use of valproic acid is associated with a high level of blood ammonia related to carnitine deficiency. This study investigates the effect of carnitine supplementation on blood ammonia levels in children with epilepsy who have been treated with valproic acid for more than six months.

Materials and Methods: This was a randomized, double-blind, placebo-controlled trial study where children with epilepsy who were treated with valproic acid were randomly allocated to the carnitine supplementation and control group. All children were followed for month, and then measured for blood ammonia level. Blood ammonia levels of both groups were compared using an Independent t-test with a significant of \( p < 0.05 \).

Results: Total of 32 children with epilepsy were enrolled as subjects in this study, with 16 children in carnitine group, and 16 children in control group. Among the subjects, 50% were male and 50% were female, with a mean age of 6.5 years old. The average duration of epilepsy in the carnitine and control group were 41.7 months and 36.9 months, respectively (\( p=0.419 \)). The duration of valproic acid therapy in the carnitine and control group were 33.1 months and 27.6 months, respectively (\( p=0.483 \)). The level of blood ammonia in carnitine and control group were 44.6 \( \mu \)g/dL and 81.4 \( \mu \)g/dL, respectively (\( p=0.007 \)).

Conclusion: The level of blood ammonia in a carnitine group was significantly lower than in a control group. It is recommended to give carnitine supplementation in epileptic children treating with long-term valproic acid.

Keywords: ammonia, carnitine, epilepsy, seizure, valproic acid
Introduction

Valproic acid is widely used in children or adults with epilepsy. This drug is very effective to control seizures in epileptic children and commonly used for a long time.\(^1,2\) The long-term used of valproic acid is associated with the increase of a serum ammonia level.\(^3\) Ammonia is toxic and causes central nervous system damaged.\(^4\) This condition should be prevented by the doctors who are treating the children with epilepsy.

Valproic acid is a short-chain fatty acid that is found for the first time in 1882 and then reported having an anti-epileptic agent in 1962.\(^1\) Until now, valproic acid is used as the first choice of anti-epileptic drug in childhood epilepsy.\(^1\) Valproic acid is used in controlling seizure as a single or combined drug in children with epilepsy. The mechanism of valproic acid in controlling seizure is by inhibiting the degradation of neurotransmitter gamma-aminobutyric acid.

The long-term used of valproic acid is associated with many side effects such as fatigue, intestinal discomfort, increased body weight, tremor, hair loss, thrombocytopenia, increased liver enzyme, and increased blood ammonia.\(^5-8\) The increased blood ammonia is a serious side effect of long-term used of valproic acid. It is associated with the liver and central nervous system damaged. The increased blood ammonia was reported to vary between 27.8% to 67.7%.\(^4,5,9\)

The high blood ammonia level related to the long-term used of valproic acid is associated with the inhibition of enzyme carbamoyl phosphate synthetase-1 and pressing the availability of acetyl-coenzyme-A that causes carnitine deficiency.\(^10\) Both of this mechanism causes the impairment of the urea cycle resulting in the accumulation of blood ammonia. The increased of blood ammonia is also caused by the disease of inborn error of metabolism, acute or chronic liver disease, acute or chronic kidney disease, Reye syndrome and using drugs 5-fluorouracil and salicylate.\(^10\)

Carnitine supplementation has been reported decreasing the blood ammonia level related valproic acid therapy for several years ago.\(^11-14\) Previously studies reported the effectiveness of carnitine supplementation in the valproic acid intoxication and psychiatric setting. However, along with our knowledge, there is no data on the effect of carnitine supplementation in epilepsy children related to valproic acid therapy. This study investigates the blood ammonia level of epileptic children treated with valproic acid for more than six months and compare its level between children who received carnitine and without carnitine supplementation.

Materials and methods

Study Design

The current study was a randomized, double-blind, placebo-controlled trial conducted between September and December 2018 at Pediatric Ward, Sanglah General Hospital, Denpasar, Bali, Indonesia. The study was approved by The Ethical Committee of the Medical Faculty, Udayana University, Sanglah General Hospital, Denpasar, Bali, Indonesia (No. 2361/UN14.2.2.VII.14/LP/2018).

Study Populations and Randomization

The study included children aged 1-18 years old who were diagnosed as epilepsy and have been treated with valproic acid for more than six months. The subjects were excluded if they suffered from the urea cycle disease, received the drugs that influenced the blood ammonia level, such as salicylate, anti-human immunodeficiency virus (HIV), and chemotherapy, and suffered from the liver and the kidney function impairment.

Subjects who met the inclusion and exclusion criteria were randomly allocated to the supplementation and control group. The random allocation was determined using a computer system with a block size of four. Both subjects and study investigators did not know the group allocation. The supplementation group received 500 mg carnitine capsule twice a day for 30 days. The control group received a similar capsule contain saccharin lactis twice a day for 30 days. The control group received a similar capsule contain saccharin lactis twice a day for 30 days.

The number of subjects were enough to be a sample, 7 subjects did not meet the inclusion criteria because they received valproic therapy for less than 6 months. A total of 38 subjects were randomly allocated, with 19 subjects in each group. On the follow-up, 6 subjects dropped out. In the end, 32 subejcts completed the study, with 16 subhects in each group.

Data Collection

Data regarding age, gender, body mass index (BMI), duration of epilepsy, duration of valproic acid therapy, serum glutamic pyruvic transaminase (SGPT), serum glutamic oxaloacetic transaminase (SGOT), blood urea nitrogen (BUN), and serum creatinine (SC) were collected.
at the beginning of the study. The children were followed and monitored for the side effects during the study period. In the end, the blood ammonia level was measured in both groups. Blood ammonia was measured using enzymatic glutamate dehydrogenase in Quantum Laboratory, Denpasar, Bali.

**Statistical Analysis**

The minimal sample size was 16 subjects each group according to the power 80% ($Z_\beta=0.842$), $\alpha=0.05$ ($Z_\alpha=1.96$). The independent t-test was done to analyzed the differences between the blood ammonia level between group. The significant level was considered if $p$-value$<0.05$. All analysis was performed using SPSS software (IBM Coorporation, Armonk, NY , USA) version 20.0.

**Results**

A total of 32 children completed the study, 16 males and 16 females, respectively. The mean aged of children in the supplementation and control group was 83.9±35.5 months and 72.2±40.6 months, respectively (Table 1). The study characteristics, such as gender, age, BMI, duration of epilepsy, duration of valproic acid therapy, SGPT, SGPT, BUN, and SC were not statistically different between groups ($p>0.05$). On the other hand, the blood ammonia level between groups at the end of the study was statistically significant with $p$-value=0.007 (Table 2).

**Discussion**

This study found that the children with epilepsy who were treated with valproic acid and received carnitine supplementation tend to have lower blood ammonia levels compared with the children without carnitine supplementation. The blood ammonia level in the carnitine supplementation group and the control group was 44.6 µg/dL and 81.4 µg/dL, respectively. It was the 45.2% lower of the blood ammonia level in the carnitine group than the control group. The result is essential and is recommended to be implemented. It could be suggested to give carnitine supplementation in children with epilepsy who have been treated with long-term valproic acid in controlling seizure.

Along with our knowledge, the current study is the first study investigating the effectiveness of carnitine supplementation on blood ammonia in children with epilepsy. Other studies on different settings reported similar results. A study on psychiatric settings reported that the carnitine supplementation doses of 30 mg/kg/day during 3 months reduced blood ammonia level from 108.3 µg/dL to 93.1 µg/dL. Unfortunately, this result was not statistically significant. Study in an adult with valproic acid intoxication (plasma level>100 µg/mL) reported that carnitine supplementation with doses of 1800 mg/day reduced the blood ammonia from 264 µmol/dL to 64.1 µmol/dL within 72 hours. They concluded that carnitine supplementation

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Supplementation Group (n=16)</th>
<th>Control Group (n=16)</th>
<th>$p$-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender, male/female</td>
<td>8/8</td>
<td>8/8</td>
<td>0.638</td>
</tr>
<tr>
<td>Age (mo.), mean±SD</td>
<td>83.9±35.5</td>
<td>72.2±40.6</td>
<td>0.394</td>
</tr>
<tr>
<td>BMI (kg/m²), mean±SD</td>
<td>17.2±3.1</td>
<td>16.2±3.7</td>
<td>0.402</td>
</tr>
<tr>
<td>Duration of epilepsy (mo.), mean±SD</td>
<td>41.7±31.3</td>
<td>36.9±24.9</td>
<td>0.419</td>
</tr>
<tr>
<td>Duration of VPA (mo.), mean±SD</td>
<td>33.1±24.1</td>
<td>27.6±18.9</td>
<td>0.483</td>
</tr>
<tr>
<td>SGOT (µ/L), mean±SD</td>
<td>27.1±2.2</td>
<td>27.1±5.7</td>
<td>0.960</td>
</tr>
<tr>
<td>SGPT (µ/L), mean±SD</td>
<td>16.1±5.2</td>
<td>14.6±4.2</td>
<td>0.403</td>
</tr>
<tr>
<td>BUN (mg/dL), mean±SD</td>
<td>21.3±3.1</td>
<td>22.3±4.5</td>
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<tr>
<td>SC (mg/dL), mean±SD</td>
<td>0.3±0.01</td>
<td>0.3±0.07</td>
<td>0.874</td>
</tr>
</tbody>
</table>

BMI: body mass index; BUN: blood urea nitrogen; SC: serum creatinine; SD: standard deviation; SGOT: serum glutamic oxaloacetic transaminase; SGPT: serum glutamic pyruvic transaminase; VPA: valproic acid.
Carnitine Supplementation on Blood Ammonia Level in Epilepsy Children

Sidiartha IGL, et al.

Table 2. The blood ammonia level in the supplementation and the control group.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Supplementation Group</th>
<th>Control Group</th>
<th>Mean Difference (95%CI)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blood ammonia, µg/dL, mean (SD)</td>
<td>44.6</td>
<td>81.4</td>
<td>36.8 (11.1 – 62.7)</td>
<td>0.007</td>
</tr>
</tbody>
</table>

CI: confidence interval; SD: standard deviation.

was effective in reducing the blood ammonia and it could be used preventing encephalopathy related to valproic acid intoxication.\(^\text{12}\)

L-carnitine has a potential capacity in treating metabolic encephalopathy caused by a high level of blood ammonia.\(^\text{13,14}\) In humans, ammonia is a product of protein and amino acid metabolism. Ammonia is a toxic substance, especially on the central nervous system and it should be degraded to urea through the urea cycle in the liver. The mechanism of the urea cycle in the liver needs several enzymes. In liver diseases, cells damaged and enzyme deficiency cause accumulation of blood ammonia resulting in hepatic hyperammonemia encephalopathy and death. Valproic acid intoxication also cause hyperammonemia encephalopathy that could be prevented with carnitine supplementation.\(^\text{10}\)

The limitation of the study has not measured the level of the blood valproic acid, so it is difficult to consider that the high level of the blood ammonia associated with the high level of blood valproic acid. Another limitation was not measuring the level of the blood carnitine. It makes difficult in measuring the increase of the blood ammonia level is caused by the decrease of the blood carnitine related to valproic acid therapy.

Conclusion

The study concludes that carnitine supplementation in children with epilepsy who were treated with long-term valproic acid was effective in lowering the blood ammonia level. It is recommended that epileptic children treating with long-term valproic acid should be added carnitine supplementation.

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References