Abstract

Objectives: Autism Spectrum Disorder is a neurodevelopmental disease with an average diagnosis age of over 3 years. Carnitine levels in ASD are important because they show potential mitochondrial dysfunction and abnormal fatty acid metabolism. In this study, in ASD children carnitine levels in dried blood spot samples were evaluated and compared with the control group.

Methods: Twenty-three children diagnosed with ASD in Research and Training Hospital (19 boys, 4 girls) and age and gender matched 24 children without ASD were enrolled in this study. 17 carnitines in dried blood samples were measured with LC-MS/MS.

Results: C0, C2, C4-OH, C5, C5-OH, C6, C16, C18 carnitines were lower (p value 0.037, 0.010, 0.005, 0.032, 0.005, 0.003, 0.043, 0.003, respectively) and C18:1 carnitine was higher (p<0.025) in ASD group compared with control group.

Conclusions: Comprehensive carnitine levels for ASD are important to establish a treatment protocol for the treatment of ASD behavior and severity. C18:1 carnitine, detected for the first time in the cases with ASD, is important for its high levels and for being a glycine transporter two inhibitor. In ASD cases, the molecular analysis might be suggested for enzymes involved in carnitine metabolism and for glycine transporter 2.

Keywords: autism spectrum disorder; C18:1 carnitine; carnitine; dried blood spot samples; liquid chromatography–tandem mass spectrometry.

Introduction

Autism spectrum disorder (ASD) is a neurodevelopmental disorder with a heterogeneous etiology and a strong genetic basis reveals itself by various clinical symptoms. The main symptoms reveal themselves as deficiencies in social communication and interaction, repetitive behavior patterns, limited interests or activities [1, 2]. Beyond the basic behavioral symptoms in individuals with ASD, gastrointestinal, immune, hepatic and endocrine systems are seen to be affected. Common comorbidities include neurological, psychiatric and physical conditions: neurological comorbidities include epilepsy, sleep disturbance, sensory abnormalities, and motor function delays and/or deficiencies; psychiatric conditions include depression, anxiety, irritability and attention deficit hyperactivity disorder, and physical health problems include chronic gastrointestinal discomfort [3]. The incidence of one or more non-ASD developmental diagnoses is as high as 83% [4]. It is associated with impaired quality of life and the lifetime cost of supporting an individual with ASD is a very high economic expense due to comorbid disorders [2, 5]. Estimates of total ASD prevalence ranged from 13.1 to 29.3 per 1,000 children over the age of 8 years. Males have four times more ASD than females [1]. ASD prevalence estimates also vary according to race and ethnicity [6].

Genetic causes of ASD are identified in approximately 30–35% of cases. For the remaining 65–70% of cases, there is a general agreement among researchers that ASD may be caused by the interaction of the combination of...
environmental and different genetic factors. Environmental factors can also take part and be reflected in epigenetic modifications [7].

Carnitine (B-hydroxy-y-N-trimethylaminobutyric acid) is necessary for the transfer of long-chain fatty acids along the internal mitochondrial membrane for the continuation of β-oxidation [8–10]. Carnitine plays a critical role in the energy balance between the cell membranes and in the energy metabolism of tissues that obtain most of their energy from fatty acid oxidation, such as cardiac and skeletal muscles. Although carnitine plays a major role in free fatty acid metabolism, carnitine increases the use of carbohydrates [11]. Carnitine can be synthesized endogenously by the human body, is very low in vegetables and grains, moderate in dairy products and high in meat, and can also be provided by the food supply. Therefore, genetic disorders in carnitine biosynthesis do not routinely cause low plasma carnitine levels [8–10, 12].

Carnitine deficiency has primary and secondary genetic forms. Secondary deficiency is caused by various fat oxidation defects and organic acidemias that lead to carnitine deficiency due to the loss of urine of acylcarnitines accumulated to enzyme deficiency. Secondary deficiency is caused by various fat oxidation defects and organic acidemias that lead to carnitine deficiency due to the loss of urine of acylcarnitines accumulated to enzyme deficiency. Primary systemic carnitine deficiency is caused by biallelic loss of function mutations in the SLC22A5 gene encoding plasma membrane organic cation transporter 2 (OCTN2) [10, 13, 14]. OCTN2 deficiency is characterized by excessive carnitine loss in the urine leading to systemic deficiency associated with skeletal myopathy, cardiomyopathy, fatty liver and hypoglycemia. Although the possibility of primary systemic carnitine deficiency due to a defect in carnitine biosynthesis has long been presumed a long time ago, primary carnitine biosynthesis disorders have not been described to date [10, 14].

Administration of carnitine to patients is the main approach of treatment for systemic carnitine deficiency and is useful in some genetic forms of secondary carnitine deficiency. Administration of carnitine and acetylcarnitine had been investigated for the treatment of many diseases including diabetic peripheral neuropathy, heart failure and mitochondrial disorders, and as an antioxidant [10].

Although autism spectrum disorder has been considered to be highly inherited and caused by many gene interactions, no single gene has been identified that adequately explains the alarmingly increasing prevalence and complex heterogeneity of the disease. As a result of these studies, the relationship between L-carnitine and autism is based on observations such as abnormal mitochondrial function occurring in ASD cases, the relationship between L-carnitine levels and the severity of autism, and genetic aspects of L-carnitine metabolism [15].

Carnitine levels in ASD are important because they show potential mitochondrial dysfunction and abnormal fatty acid metabolism [16]. Although in their studies of Filipek et al. [17], and Mostafa et al. [18], low carnitine levels in ASD children, besides, in their studies of Geier et al. [19], and Fahmy et al. [20], improvement in cases administered L-carnitine in ASD symptoms have been demonstrated, there are not many studies on this subject. For all these reasons, in this study, in ASD children carnitine levels in dried blood spot samples were evaluated and compared with the control group.

Materials and methods

The individuals who were diagnosed with ASD (DSM 5) and whose carnitine levels were studied included in the study. The results of the individuals with ASD between 2016 and 2018 were discussed. The control group consisted of the volunteer individuals who did not have any neurological and metabolic findings and have applied to the other clinics of the hospital. The study was approved by the local ethics committee (No: 05.10.2018-144).

A total of 23 children who were diagnosed ASD by Children and Adolescence Clinics of Hospital were included in the study, (19 boys aged between 5 and 18 y/o and four girls aged between 5 and 16 y/o). A total of 24 children were included in this study as a control group, including age-matched 19 boys and five girls. All participants did not have any special diets, and the group with ASD had one Pica disorder. However, the participants were not diagnosed any metabolic disorders.

The samples for fasting carnitine levels in our hospital are taken into dried blood spot card (Ministry of Health Dried Blood Spot Filter Paper for phenylketonuria screening Lot: 77/427W-112, Turkey). All samples were studied with Tandem Gold liquid chromatography-tandem mass spectrometry (LC-MS/MS) System (Zivak Technologies, Turkey) and original kits (Ref. no: ZV-3017-0500-15) used in our routine laboratory. Briefly, the test principle is that carnitines are extracted from dry blood samples by organic solvents that includes internal standard. Extracted analytes are derivatized with reagents. Derivatized carnitines are analyzed in LC-MS/MS system. Isotope dilution method is used to calculate the results. The concentration of analytes is calculated according to internal standard concentration fields. A total of 17 carnitine levels including free L-carnitine (FC or C0), acetyl-L-carnitine (ALC or C2); propionyl-L-carnitine (C3); butyryl-L-carnitine (C4); 3-hydroxybutyryl-L-carnitine (C4-OH); isovaleryl-L-carnitine (C5); glutaryl-L-carnitine (C5-DC); 3-hydroxyisovaleryl-L-carnitine (C5-OH); hexanoyl-L-carnitine (C6); adipyl-L-carnitine (C6-DC); octanoyl-L-carnitine (C8); decanoyl-L-carnitine (C10); dodecanoyl-L-carnitine (C12); myristoyl-L-carnitine (C14); palmitoyl-L-carnitine (C16); stearoyl-L-carnitine (C18); oleoyl-L-carnitine (C18:1) were measured with the device.
Statistical analysis

Statistical analyses were carried out using SPSS (Statistical Package for Social Sciences, Chicago, IL, USA) software. Shapiro-Wilk test was used for normal distribution of data and the Mann-Whitney U test was performed for the difference between groups. p<0.05 was considered statistically significant.

Results

The mean diagnosis age of children with ASD was 3.00. Comorbidities were Mild (11 children), Moderate (9 children) severe (2 children) and Profound (0 children) intellectual disability, borderline intellectual functioning (BIF) (1 child). There was a child with an EEG defect, there 2 (two) children having seizures (Table 1).

In the ASD group, 18 children were born with normal spontaneous vaginal delivery (NSVD), and five children with caesarean section (C/S); in the normal group, 14 children were born with NSVD and 10 children with C/S. Some of the ASD children have relatives and/or family members with ASD (10 children and two of them were siblings) where there was only one child in the control group. Children with ASD were using or stopped using medications (risperidone, Table 1: Clinical and family characteristics of children included in the study.

|                      | ASD (n=23) | Control (n=24) |
|----------------------|------------|---------------|
| Gender               |            |               |
| Female:4 (17.3%)     | Female: 5 (20.8%) |
| Male: 19 (82.6%)     | Male: 19 (79.1%) |
| Chronological age, year* | 11.52 (min:5.10–max:18.27) | 11.64 (min:5.43–max: 18.37) |
| Weight, kg*          | 47.61 (min:19–max:85) | 41.38 (min:18–max:78) |
| Height, cm*          | 147.22 (min:114–max:173) | 146.17 (min:110–max:176) |
| Age of diagnosis, years | 3.00 (SD: 1.23, min: 1.20–max: 5) | No |
| Comorbidity          |            |               |
| ADHD: 30.4% (n=7)    | No         |
| Pica: 4.3% (n=1)     |            |
| Stuttering: 4.3% (n=1) | No         |
| Accompanying intellectual disability |            |               |
| Mild:11              | No         |
| Moderate:9           |            |
| Severe:2             |            |
| Profound:0           |            |
| BIF: 1               |            |
| Accompanying physical illness | Visual disabilities: 2 | No |
| Hearing disability: 2 |            |
| BMI >25: 2           |            |
| No: 17               |            |
| Seizure history      |            |               |
| Yes: 2               | No         |
| No: 21               |            |
| Drug use             |            |               |
| Yes: 16              | Yes: 2     |
| No: 7                | No: 22     |
| Birth trauma         |            |               |
| Yes: 4               | Yes: 1     |
| No: 17               | No: 23     |
| Unknown: 2           |            |
| Birth type           |            |               |
| NSVD: 18             | NSVD: 14   |
| C/S: 5               | C/S: 10    |
| Age of mother (at birth) | 31.09 (min:22–max: 41) | 31.17 (min:22–max: 38) |
| Age of father (at birth) | 36.57 (min:26–max: 52) | 35.33 (min:27–max: 43) |
| Consanguineous marriage | Yes: 9     | Yes: 3      |
| No:14                | No:21      |
| The presence of autism in the family | Yes:10     | Yes: 1      |
| No: 13               | No: 23     |
| The presence of psychiatric disease in the family | Yes: 14    | Yes: 4      |
| No: 9                | No: 20     |
| Smoking in the mother during pregnancy | Yes: 10    | Yes: 2      |
| No: 13               | No: 22     |
| Smoking in the father | Yes: 11    | Yes: 16     |
| No: 12               | No: 8      |

*p value>0.05 Mann-Whitney U test. BIF, Borderline Intellectual Functioning; BMI, Body mass index; NSVD, Normal spontaneous vaginal delivery; C/S, Cesarean section.
atomoxetine, haloperidol, naltrexone, aripiprazole) for major symptoms such as behavior problems, aggression, offensiveness. In the control group, one child was using medication for anxiety disorder and one child for asthma (fluoxetine, hydroxyzine, salbutamol, montelukast and desloratadine) (Table 1).

Familial characteristics of the children were; The maternal age (at birth) was 31 (min: 22–max: 41) in the ASD group and 31 (min: 22–max: 38) in the control group; mean age of fathers (at birth) was 36 (min: 26–max: 52) in the ASD group and 31 (min: 22–max: 41) in the control group (respectively, p value 0.037, 0.010, 0.005, 0.032, 0.005, 0.003, 0.043, 0.003), and a significant decrease in C0, C2, C4-OH, C5, C5-OH, C6, C16, C18 carnitines compared to the control group (respectively, p value 0.037, 0.010, 0.005, 0.032, 0.005, 0.003, 0.043, 0.003), and a significant increase in C18:1 carnitine (p<0.025) was observed in the mean dried blood spot samples (Table 2) (Figure 1).

| Carnitine | Grup | N  | Mean, μmol/L | SD   | Reference ranges, μmol/L | Limits of detection (μmol/L) |
|-----------|------|----|--------------|------|--------------------------|-----------------------------|
| C0*       | ASD  | 23 | 35.699       | 9.299| 12–220                   | 0.023                       |
| Control   |      | 24 | 41.594       | 9.215|                          |                             |
| C2*       | ASD  | 23 | 16.365       | 3.627| 5–85                     | 0.042                       |
| Control   |      | 24 | 19.410       | 3.913|                          |                             |
| C3        | ASD  | 23 | 2.086        | 0.630| 0–6.5                    | 0.024                       |
| Control   |      | 24 | 2.441        | 0.808|                          |                             |
| C4        | ASD  | 23 | 0.276        | 0.119| 0–2                      | 0.054                       |
| Control   |      | 24 | 0.267        | 0.072|                          |                             |
| C4-OH*    | ASD  | 23 | 0.149        | 0.060| 0–0.5                    | 0.047                       |
| Control   |      | 24 | 0.175        | 0.030|                          |                             |
| C5*       | ASD  | 23 | 0.159        | 0.047| 0–1.49                   | 0.023                       |
| Control   |      | 24 | 0.191        | 0.054|                          |                             |
| C5-OH*    | ASD  | 23 | 0.316        | 0.130| 0–1.2                    | 0.022                       |
| Control   |      | 24 | 0.417        | 0.139|                          |                             |
| C5-DC     | ASD  | 23 | 0.194        | 0.060| 0–0.35                   | 0.076                       |
| Control   |      | 24 | 0.217        | 0.054|                          |                             |
| C6*       | ASD  | 23 | 0.092        | 0.028| 0–0.7                    | 0.041                       |
| Control   |      | 24 | 0.114        | 0.020|                          |                             |
| C6-DC     | ASD  | 23 | 0.087        | 0.041| 0–0.5                    | 0.024                       |
| Control   |      | 24 | 0.087        | 0.031|                          |                             |
| C8        | ASD  | 23 | 0.140        | 0.038| 0–0.9                    | 0.088                       |
| Control   |      | 24 | 0.130        | 0.030|                          |                             |
| C10       | ASD  | 23 | 0.177        | 0.064| 0–0.7                    | 0.088                       |
| Control   |      | 24 | 0.161        | 0.044|                          |                             |
| C12       | ASD  | 23 | 0.140        | 0.043| 0–1.44                   | 0.068                       |
| Control   |      | 24 | 0.140        | 0.030|                          |                             |
| C14       | ASD  | 23 | 0.139        | 0.036| 0–1.1                    | 0.067                       |
| Control   |      | 24 | 0.139        | 0.033|                          |                             |
| C16*      | ASD  | 23 | 1.123        | 0.277| 0–2.1                    | 0.077                       |
| Control   |      | 24 | 1.380        | 0.437|                          |                             |
| C18*      | ASD  | 23 | 0.839        | 0.213| 0–7                      | 0.042                       |
| Control   |      | 24 | 1.230        | 0.470|                          |                             |
| C18:1*    | ASD  | 23 | 0.547        | 0.115| 0–4                      | 0.091                       |
| Control   |      | 24 | 0.478        | 0.151|                          |                             |

SD, Standard deviation. *p value<0.05 significant.
Figure 1: Carnitines with statistical significant difference between ASD and control group.
Discussion

Carnitine levels were analyzed with dried blood spot samples by LC-MS/MS method in 23 ASD cases. According to the control group, carnitines of C0, C2, C4-OH, C5, C5-OH, C6, C16 and C18 are low and carnitine of C18:1 is high, other carnitine levels were similar.

In their meta-analysis of Rossignol and Frye [21], it has been reported that the prevalence of mitochondrial disorders (5%) was significantly higher in cases with ASD than in the general population and that ASD was associated with mitochondrial disorders. The prevalence of the biomarker values of mitochondrial dysfunction with ASD was much higher than the prevalence of mitochondrial disorders. It has been reported that the values of many mitochondrial biomarkers (lactate, pyruvate, carnitine, and ubiquinone) differ significantly between ASD and controls, and some markers are associated with the severity of ASD [21]. Carnitine is thought to exert protective effects on mitochondria and all cells by inhibiting free fatty acid-induced mitochondrial membrane damage and/or secondary effects [22, 23].

In our study, it has been reported that for certain symptoms drugs were administered by some of the ASD children or some of them stopped the treatment, in one of the cases cortexin (contains polypeptides, vitamins, minerals and amino acids) was administered but was quit for a long time ago, in two cases probiotics and dietary supplements (fish oil and B vitamins) were received. In our study, there were no cases using carnitine. Among ASD cases with mitochondrial disorders, the use of carnitine is highlighted as one of the most recommended nutritional supplements [21, 24, 25].

In the study of Geier et al. [19], it has been reported that there is a significant improvement in several clinical measurements of ASD severity, that is in 30 cases diagnosed with ASD and randomly assigned as liquid L-carnitine (n=19) and placebo (n=11) for 3 months. Similarly, Fahmy et al. [18] have been reported that 30 cases diagnosed with autism were randomly assigned to receive of liquid l-carnitine (n=16) or placebo (n=14) for 6 months, and after 6 months, liquid l-carnitine therapy administered significantly improved the autistic behavior and decreased the severity of autism. Total and free carnitine levels were examined in the studies [19, 20]. Based on the data obtained from these studies and our study, it is obvious that carnitine levels are important for ASD and it should be emphasized that carnitine levels should be examined in terms of ASD treatment protocols.

In the study of Lv et al. [16], it has been reported that levels of free carnitine (C0), glutaryl carnitine (C5-DC), octyl carnitine, carnosyl carnitine (C26) were found to be significantly lower in 60 children (49 boys and 11 girls) diagnosed with ASD (42.86 ± 11.03 months) and 30 children (25 boys and five girls) with typically developing (39.32 ± 12.88 months). As a conclusion, it has been reported that glutaryl carnitine and carnosyl carnitine might be potential biomarkers for diagnosis of ASD, that the limitations of the study were the relatively small sample size and participants which were the children within 6 years of age [16]. In contrast to this aforementioned study, in our study, there is not any significant difference in C0, C5-DC levels compared to the control group and octyl, C26 carnitine levels were not examined. In our study, the age of the ASD group reached up to 18.27 years and it has been stated that in the ASD group compared to the control group C0, C2, C4-OH, C5, C5-OH, C6, C16, C18 levels were significantly low, C18:1 level significantly high.

Adams et al. [26] demonstrated that carnitine results in 67 cases with ASD in both groups with and without carnitine administration started at levels similar to neurotypical controls, and that treatment led to a significant increase in L-carnitine and non-significant increase in acetyl-L-carnitine. In this study, in cases with ASD, it has been stated that administered L-carnitine can be absorbed better than acetyl-L-carnitine in other studies [19, 20] and therefore might be more effective. They also reported that the benefit from carnitine was very limited in cases with ASD [26]. In this study [26] carnitine levels were initially similar to the control group so that it is considered to be the reason for the limitation in benefit from carnitine in cases with ASD. Although no protocol for treatment was established in our study, it could be said that carnitine supplementation would be beneficial in cases with ASD with low carnitine levels considering the studies in the literature [19, 20, 26]. But carnitine treatment might be harmful in cases of long-chain fatty acid oxidation disorders (FAODs), as it increases the concentrations of long-chain acylcarnitines, which are potentially arrhythmogenic [27]. Therefore, in carnitine supplementation, long-chain fatty acid oxidation disorders and side effects of carnitine should be taken into consideration.

In our study, it has been proved that only carnitine C18: 1 in the ASD group was statistically significant (p<0.025). Height C18:1 carnitine levels were also determined in carnitine palmitoyltransferase-2 [28], carnitine-acylcarnitine translocase [29], long-chain acyl-CoA dehydrogenase deficiencies [30].
Clark-Taylor and Clark-Taylor have been stated that C10:1, C14:2 and C14:1 for plasma acyl-carnitine profile were high and that long-chain acyl-CoA dehydrogenase deficiency might be a cause of autism, and it might be important to identify acyl-carnitine profiles in other children with autism [31]. We found that a high level of C18:1 suggests that the studies with long-chain acyl-CoA dehydrogenase and enzymes related to carnitine metabolism should be performed in ASD cases.

Also, carnitine C18:1 has a role as a glycine transporter 2 (GlyT2) inhibitor [32, 33]. The main role of GlyT2 is to recapture glycine released in the synaptic cleft and maintain high glycine concentration in the presynaptic neuron [34, 35]. It is considered that C18:1 carnitine might be interested in symptoms of ASD through GlyT2. Although there is a study on glycine transporter one in cases with ASD [36], no studies on GlyT2 have been found in the literature. GlyT2 inhibitors show potential as analgesics [33] and oleoyl-L-carnitine may have the potential for new research into pain sensation in ASD individuals.

The wideness of age distribution of the cases in this study is important. One of the limitations of our study is the relatively inadequate number of cases. Besides, no treatment protocol was applied in the study. Therefore, the efficacy of carnitine treatment could not be monitored. It is another limitation that the uncertainty of relationships of carnitine C18:1 with long-chain fatty acid oxidation or enzyme disorders in carnitine metabolism is not clarified as urine organic acids were not studied. In addition, this study was conducted on dry blood samples can be stated as a limitation. It might be confirmed by studies with carnitines to be measured in direct blood samples. Despite all these, it is thought that this study will contribute to the literature considering the small number of studies on carnitine levels in ASD cases.

Comprehensive carnitine levels for ASD are important for the diagnosis of ASD and the treatment in the elimination of ASD behaviors and severity. It is important to determine the relationship between ASD behavior and the severity of educational interventions and carnitine levels in cases with ASD. In our study, although the number of patients was relatively low in cases with ASD, a molecular level analysis could be recommended for enzymes playing a role in carnitine metabolism related to C18:1 high levels of carnitine and glycine transporter 2.

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