



Screening Inherited Metabolic Disorder in Children with Intellectual Disability and Epilepsy

Zeka Geriliği ve Epilepsisi Olan Çocuklarda Kalıtsal Metabolik Hastalık Taraması

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Abstract

Objective: To indicate the benefits of the screening of inherited metabolic disorders in patients with epilepsy, global developmental delay, and intellectual disability.

Materials and Methods: The medical records of 1100 patients who were investigated for inherited metabolic disorders between March 2014 and June 2017 were evaluated. Five hundred patients with epilepsy and global developmental delay/intellectual disability with mild/moderate and non-specific neurologic findings were enrolled in the study.

Results: Inherited metabolic disorders were detected in 7 of 500 patients (1.4%) with epilepsy and global developmental delay/intellectual disability. One patient was diagnosed as having tyrosinemia type-2, one had Menkes disease, one had mitochondrial disease, one had hyperphenylalaninemia, two siblings were diagnosed as having 3-methylcrotonyl Coa carboxylase deficiency, and one patient was diagnosed as having phenylketonuria.

Conclusion: The prevalence of inherited metabolic disorders is higher in countries with a high consanguinity ratio such as Turkey. Lack of the regular screening in patients with mild/moderate and non-specific neurologic findings result in late diagnosis.

Keywords: Inherited metabolic disorder, epilepsy, intellectual disability, metabolic screening

Öz

Amaç: Bu çalışmanın amacı epilepsi, yaygın gelişimsel gecikme ve zeka geriliği olan hastalarda kalıtsal metabolik hastalık taraması yapmanın yararlarını göstermektir.

Gereç ve Yöntem: Mart 2014 ve Haziran 2017 arasında kalıtsal metabolik hastalık nedeniyle araştırılan 1100 hastanın tıbbi kayıtları değerlendirildi. Çalışmaya hafif/orta ve non-spesifik nörolojik bulgusu olan 500 epilepsi ve yaygın gelişimsel gecikme/zeka geriliği olan hasta alındı.

Bulgular: Epilepsi ve yaygın gelişimsel gecikme/zeka geriliği olan 500 hastanın 7'sinde (%1,4) kalıtsal metabolik hastalık tespit edildi. Bir hastaya tirozinemi tip-2, bir hastaya Menkes hastalığı; bir hastaya mitokondriyal hastalık, bir hastaya hiperfenilalaninemi, 2 kardeşe 3-metilcrotonil CoA karboksilaz eksikliği ve bir hastaya da fenilketonüri tanısı konuldu.

Sonuç: Türkiye gibi yüksek akrabalık oranına sahip ülkelerde kalıtsal metabolik hastalık prevalansı daha yüksektir. Hafif/orta ve non-spesifik nörolojik bulguları olan hastalarda düzenli bir taramanın olmaması geç tanı ile sonuçlanır.

Anahtar Kelimeler: Kalıtsal metabolik hastalık, epilepsi, zeka geriliği, metabolik tarama

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Introduction

Inherited metabolic disorders (IMDs) are not common causes of epilepsy and intellectual disability (ID). IMDs should be considered in the presence of neonatal seizures, neonatal or infantile deaths with unknown etiology, affected family members, parental consanguinity, macrocephaly or microcephaly, hydrops fetalis, progressive global psychomotor retardation, multisystem involvement such as cardiomyopathy, hepatosplenomegaly, cataract and muscle weakness, and characteristic magnetic resonance imaging findings such as cerebral and/or cerebellar atrophy, abnormal myelination and striatal necrosis (1,2,3,4). Seizures may be a part of a more complex neurologic presentation or sometimes only a feature of IMDs. Some IMDs (40-60%) can lead to isolated or recurrent convulsions (3).

ID, which is also known as mental retardation, is characterized by impairment of cognitive functions, adaptive behavior and life skills with limitations of learning, and presentation before age 18 years. In addition, children younger than 6 years are considered to have global developmental delay (GDD) if they perform more than 2 SDS below age-matched peers. GDD affects 1-3% of children aged under 6 years, and there are 2-3% of patients with ID in different populations (5,6). Global psychomotor retardation without other potential explanations may be suggestive of an IMD. In the absence of clues for other common causes of GDD/ID, the study of metabolic tests is recommended. Focused or sequential metabolic testing can increase the diagnostic rates up to 14% (7).

Treatments of some IMDs include dietary therapy, essential amino acid and vitamin supplementation, substrate inhibition and reduction therapy, enzyme replacement therapy, hematopoietic stem cell transplantation, and gene therapy. Therefore, early recognition is very important. In addition, identification of non-treatable causes is beneficial to the individual's family and allows genetic counseling (8).

The aim of this study was to explore the benefits of screening for IMDs in patients with epilepsy and GDD/ID with mild/moderate and non-specific neurologic findings.

Materials and Methods

This was a retrospective single-center clinical study. The medical records of 1100 patients who were investigated for IMDs between March 2014 and June 2017 were evaluated. Detailed disease histories, age, sex, and physical examinations were recorded in a form.

Inclusion criteria: Five hundred patients with epilepsy and GDD/ID with mild/moderate and non-specific neurologic findings were enrolled in the study.

Exclusion criteria: Patients with a specific genetic syndrome or neurologic disease, central nervous system (CNS) infection or CNS tumor, and febrile convulsions were excluded from the study. Although multidrug-resistant convulsions, cerebral palsy, and infantile severe hypotonia indicate metabolic disorders, they were also excluded from the study because the aim of this study was to explore IMDs in epilepsy and GDD/ID with mild/moderate neurologic symptoms.

Laboratory investigations: Non-specific tests include complete blood count, electrolytes, glucose levels, liver transaminases, urea, creatinine, creatine phosphokinase, thyroid function tests, vitamin B12, serum ammonia, lactate, pyruvate levels and peripheral chromosome analysis. Specific metabolic tests consist of plasma and urine amino acid analysis, blood acyl carnitine analysis, biotinidase activity and urinary organic acid analysis. Urinary glycosaminoglycan levels, urine sulfide oxide, serum copper, ceruloplasmin and homocysteine levels were studied in selected cases. All molecular genetic analyses were performed using next-generation sequencing in patients diagnosed as having IMDs.

The study was approved by the Ethics Committee of Erciyes University, Medical Faculty (Protocol number: 2017/421).

Informed consent was given by all parents of children with IMDs.

Statistical Analysis

The IBM SPSS Statistics 24.0 statistical package program was used in the evaluation of the data. Parameters with abnormal distribution are expressed as median (25th percentile 75th percentile). Data are presented as counts, percentages, and minimum and maximum values.

Results

Five hundred patients with epilepsy and GDD/ID whose physical examinations and metabolic investigations were complete, and who met the inclusion criteria of the study were enrolled. Among the 500 children, 293 were male and 207 were female. The sex ratio was 1.4/1 in favor of the males. The ages of patients ranged from 20 days to 18 years. One hundred sixty-five patients (33%) were evaluated for GDD/ID of unknown etiology. Two hundred sixty-eight patients (53.6%) were followed up for epilepsy, and 67 patients (13.4%) with epilepsy and GDD/ID. Three hundred three (90.4%) of the 335 patients with epilepsy were using single antiepileptic drugs, 32 (9.6%) were using two antiepileptic drugs. Denver Developmental Screening Test II records of 108 patients and intelligence tests of 84 patients were available. Intelligence quotient (IQ) of 50-70 was considered mild impairment, and an IQ of less than 50 was considered moderate-to-severe impairment. The characteristics of children investigated for IMDs are shown in Table 1.

IMDs were diagnosed in seven patients as follows: one patient was diagnosed as having Tyrosinemia type-2, one had Menkes disease, one had Mitochondrial disease, one had Hyperphenylalaninemia, two siblings were diagnosed as having 3-methylcrotonyl CoA carboxylase deficiency, and one had Phenylketonuria. The demographic and clinical findings of patients with IMDs are shown in Table 2 and the laboratory findings and prognoses of patients with IMDs are shown in Table 3. In this study, the prevalence of IMDs among patients with epilepsy and GDD/ID was 1.4%.

Discussion

Seizures may be a part of a more complex neurologic presentation or sometimes only feature as IMDs. Mercimek-Mahmutoglu et al. (9) evaluated 150 patients who underwent

lumbar puncture due to epilepsy and movement disorder. IMDs were diagnosed in 6 (4%) of 150 patients. Sixty-six (44%) of 150 patients had GDD and epilepsy. IMDs were found in 1/268 (0.37%) of patients with epilepsy and 2/67 (2.98%) of patients with epilepsy and GDD/ID in our study, similar to these findings.

ID is a heavy burden both on the individual and society because its effects continue for a lifetime and also patients with ID have increased morbidity and mortality (10). Therefore, it is very important to identify treatable IMDs. There are few studies regarding ID and IMDs. Treatable IMDs were identified in 1-5% of patients as the cause of ID in some reports (11,12). Treatable IMDs, which have been detected in our study, are phenylketonuria, tyrosinemia type 2, mitochondrial disorders,

3-methylcrotonyl CoA carboxylase deficiency, and Menkes disease.

Engbers et al. (13) evaluated 433 patients with neurodevelopmental disorders and showed that 3% of these patients had IMDs. IMDs were found in 4/165 (2.4%) of patients with GDD/ID in our study.

Papavasiliou et al. (14) evaluated 118 patients with unexplained developmental delay ages from 3 months to 13 years and found 16 (13.6%) patients with neurometabolic disorder. The higher rate of IMDs in this report was attributed to more specific testing being performed such as cerebrospinal fluid lactate, very long-chain fatty acids, and mitochondrial enzymes.

van Karnebeek and Stockler-Ipsiroglu (15) reported that non-targeting screening with plasma amino acids, total homocysteine, acyl carnitine profile, copper, ceruloplasmin, urine organic acid, purine and pyrimidines, creatine metabolites, oligosaccharides, and glycosaminoglycans identified 64% of treatable IMDs. Diagnosis of all our patients was performed using non-targeting screening. The blood acyl carnitine profile and amino acid analysis are very important in the first step of specific metabolic investigations, and LC/MS/MS analysis allows the diagnosis of a large number of IMDs alone.

Henderson et al. (16) screened 1087 patients with mental retardation and found phenylketonuria (PKU) in three patients, cystinuria in two patients, and Hartnup disease in one patient. The overall frequency of IMDs was found as 0.6%. In our study, the frequency of IMDs among patients with epilepsy and GDD/ID was 1.4%.

PKU is a well-known cause of ID. Papassin et al. (17) reported PKU in a patient with ID at 20 years of age, like our case. The frequency of PKU was reported as 1/1581 in 4744 children with mental retardation (18). The PKU frequency was 1/165 in patient with unexplained GDD/ID in our study.

The incidence of IMDs is remarkably high in the Turkish population, which is partially due to the high rate of consanguinity. A total of 572 cases with 12 different types of aminoacidopathies were detected in 10,800 Turkish children with ID (19). PKU (4.7%) and homocystinuria (0.2%) were common causes of ID in this report. The detection of Hyperphenylalaninemia in one of seven patients diagnosed with IMD and the detection of PKU in one patient supports this study.

Table 1. The characteristics of the children investigated for inherited metabolic disorders

Variables	Total number of patients	(%)
Variables	500	100
Sex distribution		
Male	293	58.6
Female	207	41.4
GDD/ID	165	33
Epilepsy	268	53.6
Epilepsy and GDD/ID	67	13.4
DDST II (n=108)		
Language development	19	17.60
Gross motor development	38	35.19
Retardation in two areas	12	11.11
Retardation in three areas	13	12.03
Retardation in all areas	26	24.07
Intelligence test score (n=84)		
Borderline intelligence	7	8.33
Mild mental retardation	49	58.33
Moderate mental retardation	28	33.34

DDST II: Denver Developmental Screening Test II, GDD: Global developmental delay, ID: Intellectual disability

Table 2. The demographic and clinical findings of the patients with inherited metabolic disorders

Case	Age	Sex	Consanguinity	Clinical findings
1	8 months	M	(-)	Epilepsy, poor head control and hair changes
2	4 years	M	(+)	Complex partial seizure, mild ID in Stanford-Binet Intelligence Scale and intermittent photophobia
3	10 years 4 months	F	(-)	Headache, learning disability, attention deficit hyperactivity disorder (using methylphenidate) and mild ID in WISC-R
4	9 years	M	(+)	Generalized tonic clonic seizures (using valproic acid) and mildly impaired concentration difficulty in recent months.
5	17 years	M	(+)	Poor school performance and mild ID in WISC-R
6	9 years 2 months	F	(+)	Convulsion and mild ID in WISC-R
7	15 years	M	(+)	Learning disability and mild ID in the WISC-R

WISC-R: Wechsler Intelligence Scale for Children, ID: Intellectual disability

Table 3. The laboratory findings and prognosis of the patients with inherited metabolic disorders

Case	Laboratory findings	Cranial MRI findings	Mutation	Diagnosis	Prognosis
1	Pili torti, low serum ceruloplasmin and copper level*	Diffuse cerebral and cerebellar atrophy in MRI, vascular tortuosity in middle cerebral and vertebrobasilar arteries in cranial MRA	c.3352G>A in ATP 7A gene	Menkes disease	He is bedridden and he has a significant feeding difficulty
2	Elevated plasma tyrosine and N-acetyl tyrosine level in urinary organic acids analysis†	Normal	c.935T>C and c.1223C>T in TAT gene	Tyrosinemia type 2	Seizures were under control after dietary therapy
3	Basic metabolic tests were normal	Hyperintense signal changes in the cerebellum close to the dentate nucleus and subcortical deep white matter in the cerebral hemispheres	m.8860A>G in ATP6 gene	Mitochondrial disease	Behavioral disorders improved partially with coenzyme Q10, carnitine and vitamins
4	Plasma phenylalanine level was 524 µmol/L	Normal	Not performed	HFA (not responsive to a BH4 loading test)	Seizures were controlled after phenylalanine restricted diet
5	Plasma phenylalanine level was 1836 µmol/L	Hyperintense areas adjacent to posterior horn of both lateral ventricles	c.782G>A in PAH gene	PKU (not responsive to a BH4 loading test)	School performance is bad and is not fully following the diet
6	C5-OH level elevated in LC/MS/MS analysis‡	Normal	Not performed	3-methylcrotonyl CoA carboxylase deficiency	No improvement in behavior and recognition with leucine restricted diet and carnitine treatment.
7	C5-OH level elevated in LC/MS/MS analysis§	Normal	Not performed	3-methylcrotonyl CoA carboxylase deficiency	School performance is not so bad (only using carnitine)

*Serum ceruloplasmin concentration was 0.08 g/L (normal range: 0.31-0.91) and serum copper was 22 µg/dL (normal range: 50-155).
†Plasma tyrosine concentration was 1450 µmol/L (normal range: 0-400) in plasma amino acid analysis. 4-OH phenyl acetic acid concentration was 915 mmol/molCre (normal range: 6-24), 4-OH phenyl lactic acid was 9298 mmol/molCre (normal range: 0-2) N-acetyl tyrosine concentration was 719 mmol/molCre (normal range: 0-2) in urinary organic acids analysis.
‡C5-OH concentration was 9 µmol/L (normal range: 0-1.2) in LC-MS analysis. 3-methylcrotonyl glycine level was 588.9 mmol/molCre (n<2), 3-hydroxyisovaleric acid concentration was 1410.5 mmol/molCre (n=0-46) and 4-hydroxyphenylacetic acid concentration was 1020 mmol/molCre (n=6-28) in urinary organic acid analysis.
§C5-OH concentration was 24.8 µmol/L (normal range: 0-1.2) in LC-MS analysis. 3-methylcrotonyl glycine concentration was 874 mmol/molCre (n<2), 3-hydroxyisovaleric acid value was 1778 mmol/molCre (n=0-46) in urinary organic acid analysis.
MRI: Magnetic resonance imaging, MRA: Magnetic resonance angiography, HFA: Hyperphenylalaninemia, PKU: Phenylketonuria

Sempere et al. (20) evaluated 944 patients with unexplained ID and reported three patients with cerebral creatine deficiency syndromes, one patient with adenylosuccinate lyase deficiency, and three patients with PKU.

Study Limitations

The limitation of this study is that it did not allow screening of creatine biosynthesis, gamma-aminobutyric acid catabolism, purine and pyrimidine metabolism, congenital glycosylation, and glucose transport defects, which can present with nonspecific ID.

Conclusion

Early diagnosis and treatment is very important, and also identification of non-treatable causes is beneficial to the affected

individual's family and allows genetic counseling. The prevalence of IMDs is higher in countries with a high consanguinity ratio such as Turkey. The lack of regular screening in patients with mild/moderate and non-specific neurologic findings result in late diagnosis.

Ethics

Ethics Committee Approval: The study was approved by the Ethics Committee of Erciyes University, Medical Faculty, (Protocol number: 2017/421).

Informed Consent: Consent form was filled out by all parents of cases diagnosed with inherited metabolic disorders.

Peer-review: Internally peer-reviewed.

Authorship Contributions

Surgical and Medical Practices: P.S.Ü., A.S.G., H.G.P., Concept: P.S.Ü., Design: P.S.Ü., A.S.G., Y.A.T., Data Collection

or Processing: H.G.P., İ.G., S.G., Analysis or Interpretation: M.K., F.K., Literature Search: P.S.Ü., İ.G., Writing: P.S.Ü.

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