MCT oil-based diet reverses hypertrophic cardiomyopathy in a patient with very long chain acyl-coA dehydrogenase deficiency

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Very long chain acyl-CoA dehydrogenase (VLCAD) deficiency is one of the genetic defects of mitochondrial fatty acid beta-oxidation presenting in early infancy or childhood. If undiagnosed and untreated, VLCAD deficiency may be fatal, secondary to cardiac involvement. We assessed the effect of replacing part of the fat in the diet of a 2 ½-month-old male infant, who was diagnosed with VLCAD deficiency, with medium-chain triglyceride (MCT) oil and essential fats. The patient presented with vomiting, dehydration, and was found to have persistent elevation of liver function tests, hepatomegaly, pericardial and pleural effusion, right bundle branch block, and biventricular hypertrophy. Because of the cardiomyopathy, hepatomegaly, and an abnormal acylcarnitine profile and urine organic acids, he was suspected of having VLCAD deficiency. This was confirmed on acyl-CoA dehydrogenase, very long chain (ACADVL) gene analysis. He was begun on an MCT oil-based formula with added essential fatty acids, uncooked cornstarch (around 1 year of age), and frequent feeds. By 7 months of age, cardiomyopathy had reversed and by 18 months of age, all cardiac medications were discontinued and hypotonia had improved such that physical therapy was no longer required. At 5 years of age, he is at the 50th percentile for height and weight along with normal development. Pediatricians need to be aware about the basic pathophysiology of the disease and the rationale behind its treatment as more patients are being diagnosed because of expansion of newborn screen. The use of MCT oil as a medical intervention for treatment of VLCAD deficiency remains controversial mostly because of lack of clear phenotype-genotype correlations, secondary to the genetic heterogeneity of the mutations. Our case demonstrated the medical necessity of MCT oil-based nutritional intervention and the need for the further research for the development of specific guidelines to improve the care of these patients.

Key words: Cardiomyopathy, carnitine, inborn error of metabolism, MCT oil, very long chain acyl-CoA dehydrogenase deficiency.

Introduction

Very long chain acyl-CoA dehydrogenase (VLCAD) is the first enzyme in the beta oxidation of fatty acids once they have been transported inside the mitochondria;¹² the enzyme lies on the inner mitochondrial membrane. During times of high energy requirements, long chain fatty acids are mobilized from adipose tissue and eventually metabolized by the heart and muscle. There are three phenotypes associated with this enzyme deficiency based on age of presentation and organ system involvement.³ In this case report, we describe a patient with VLCAD deficiency diagnosed symptomatically after he was found to have hepatomegaly and cardiomyopathy.

Case Report

Our patient was a 2½-month-old male infant who presented to the hospital because of persistent emesis for 24 hours. He was also found to be hyponatremic, dehydrated, and hypotensive. The patient was admitted...
Table 1: Acylcarnitine profile showing marked elevation of long chain species, especially C14 complex, C16, C18:1 in a pattern consistent with VLCAD deficiency

| Species | Acyl Group | Normal Range | Result | Status |
|---------|------------|--------------|--------|--------|
| C2      | Acetyl     | 2.0-15.7     | 0.81   | Low    |
| C3:1    | Propenoyl  | <0.03        | BQL    | NL     |
| C3      | Propionyl  | 0.75         | BQL    | NL     |
| C4      | Butyryl/Isobutyryl | <0.43 | BQL    | NL     |
| C5:1    | Tiglyl     | 0.03         | 0.07   | Elevated |
| C5      | Isovaleryl | >0.37        | 0.05   | NL     |
| C4-OM   | 3-OH-Butyl | <0.21        | BQL    | NL     |
| C6      | Hexanoyl   | <0.25        | BQL    | NL     |
| C5-OM   | 3-OH-Isovaleryl | <0.08 | BQL    | NL     |
| B:DC    | Benzyoyl   | >0.03        | BQL    | NL     |
| C4:DC   | Methylmalonyl | 0.04 | BQL    | NL     |
| C8:1    | Octenoyl   | >0.52        | 0.06   | NL     |
| C8      | Octanoyl   | >0.22        | 0.03   | NL     |
| C5:DC   | Glutaryl   | >0.03        | BQL    | NL     |
| C6-DC   | Adipoyl    | >0.08        | 0.07   | NL     |
| C10:1   | cis-4-Decenoyl | >0.30 | 0.06   | NL     |
| C10     | Decanoyl   | >0.34        | 0.12   | NL     |
| C9:DC   | Subenyl    | >0.08        | 0.03   | NL     |
| C12-1   | Dodecenoyl | >0.24        | 0.13   | NL     |
| C12     | Dodecanoyl | >0.17        | 0.25   | Elevated |
| C14:2   | Tetradecanoyl | >0.15 | 0.73   | Elevated |
| C14:1   | Tetradecenoyl | >0.26 | 3.18   | Elevated |
| C14     | Tetradecanoyl | >0.10 | 2.02   | Elevated |
| C14:1-OM| 3-OH-C14:1 | >0.05        | 0.17   | Elevated |
| C14-OM  | 3-OH-C14   | >0.03        | 0.06   | Elevated |
| C16     | Palmitoyl  | >0.27        | 4.89   | Elevated |
| C16-OM  | 3-OH-Palmitoyl | >0.03 | 0.06   | Elevated |
| C18:2   | Linoleoyl  | >0.04        | 1.68   | Elevated |
| C18:1   | Oleoyl     | >0.42        | 6.88   | Elevated |
| C18:2-OM| 3-OH-Linoleoyl | >0.03 | 0.04   | Elevated |
| C18:1-OM| 3-OH-Oleoyl | >0.03        | 0.06   | Elevated |
| C16:DC  | C16-Dicarboxylic | >0.03 | 0.07   | Elevated |
| C18:1-DC| C18:1-Dicarboxylic | >0.03 | 0.10   | Elevated |

BQL - Below quantitation limits; NL - Normal

Discussion

VLCAD deficiency can present in one of the following three forms: (1) severe, early-onset with cardiac and multiorgan failure VLCAD-C,[6,7] (2) early childhood...
form with hypoketotic hypoglycemia and hepatomegaly VLCAD-H, and (3) the later-onset episodic myopathic form VLCAD-M. Clinically, our patient fits the first phenotype. He had a typical presentation consisting of hepatomegaly, cardiomyopathy, and cardiac arrhythmias. We did not record any hypoglycemic episodes during his hospital stay, though he was on a dextrose drip as part of his rehydration.

Treatment is essentially dietary modification, with avoidance of long-chain fatty acids and supplementation with medium chain triglycerides, so that the enzyme-deficient step can be bypassed. Such treatment should reverse most symptoms, although during times of stress, like exercise, the MCT dosage may need to be raised to supply the extra energy. This helps prevent mobilization of long-chain fatty acids from the adipose tissue. Fasting needs to be avoided for similar reasons. Close follow-up with a metabolic specialist and nutritionist is important to optimize dietary management. Acute rhabdomyolysis may occur with any form of VLCAD deficiency and must be treated aggressively because of the danger of renal failure. Also, CPK along with CK-MB (creatinine phosphokinase, muscle and brain fractions) need to be monitored. Any siblings of the patient should be evaluated, as the symptoms may be subtle.

In our clinic, an MCT oil-based diet is the main therapy for all patients with VLCAD deficiency. This treatment is based on our understanding of the pathophysiology of the condition. A recent DELPHI protocol recommends treatment based on VLCAD type and age. There was no consensus on the treatment for asymptomatic VLCAD-C patients who are being breast fed and are less than 12 months of age. A European consensus report similarly could not come to a conclusion. Patient with symptomatic VLCAD-C deficiency should have maximal MCT oil-based nutrition in both of these recommendations. The use of MCT oil as a medical intervention VLCAD deficiency remains controversial mostly because of lack of clear phenotype-genotype correlations, secondary to the genetic heterogeneity of the mutations. Though genotype-phenotype correlation have been reported in the past, these are based on patients who were diagnosed symptomatically.

Because of the expanded newborn screen and its implementation in parts of the world, more pediatricians are going to be involved in the care of these patients, and this report will help them understand the basis of treatment for these patients. We feel that once we have genotype-phenotype information available on patients diagnosed through newborn screen, we will be able to improve the treatment further and provide personalized interventions. Based on such information, further clinical studies with a larger sample size will help determine optimal nutrition care for patients with VLCAD deficiency, as well as the role of MCT oil supplementation.

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