Specialist Pembe SOYLU ÜSTKOYUNCU

Pediatric Nutrition and Metabolism

Dr. Pembe SOYLU USTKOYUNCU

Department: Pediatric Nutrition and Metabolism

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Clinic: Pediatric Nutrition and Metabolism-2

Medical Interests and Expertise:

Phenylketonuria

Organic Acidemias

Disorders of Amino Acid Metabolism

Mitochondrial Myopathies

Lysosomal Storage Diseases

Newborn Screening Tests

Other Rare Metabolic Diseases

Malnutrition

Obesity

Education:

2004-Cumhuriyet University, Medical Faculty

2010-Erciyes University, Medical Faculty, Department of Pediatrics.

2015-Erciyes University, Medical Faculty, Department of Pediatrics, Division of Pediatric Nutrition and Metabolism

Institutions:

2004-2005 Yozgat/Akdagmadeni Bozhuyuk Health Center

2005-2010 Erciyes University, Medical Faculty, Department of Pediatrics

2010-2011 Isparta/Yalvac Public Hospital

2011-2015 Erciyes University, Medical Faculty, Department of Pediatrics, Division of Pediatric Nutrition and Metabolism

2015-2018 Kayseri Training and Research Hospital, Pediatric Nutrition and Metabolism Clinic

2018-2019 Kayseri City Hospital, Pediatric Nutrition and Metabolism Clinic

Publications:

1. Dursun I, Poyrazoğlu H, Soylu P, Yılmaz A, Gürgöze MK, Gündüz Z. The Evaluation of Children with Acute Renal Failure in Pediatric Intensive Care Unit. Erciyes Medical Journal 2009;31(3):231-236.

2. Canpolat M, Poyrazoğlu HG, Yıkılmaz A, Soylu P, Per H, Gümüş H, Kumandaş S. A case of Dyke-Davidoff-Masson syndrome. Gaziantep Med J 2012;18(2): 122-124.

3. Soylu P, Kardas F, Kendirci M. Vitamin D Intoxication in Children. Turkiye Klinikleri J Pediatr Sci. 2012;8(2):148-155.

4.Kardas F, Soylu P, Kendirci M. Epidemiology of Fabry Disease. Turkiye Klinikleri J Pediatr. 2012;21(4 Suppl 1):S 7-9.

5.Gokay S, Kendirci M, Ustkoyuncu PS, Kardas F, Bayram AK, Per H, Poyrazoğlu HG. Tyrosinemia type II: Novel mutations in TAT in a boy with unusual presentation. Pediatr Int. 2016;58(10):1069-1072.

6. Gokay S, Ustkoyuncu PS, Kardas F, Kendirci M. The outcome of seven patients with hereditary tyrosinemia type 1. J Pediatr Endocrinol Metab. 2016;29(10):1151-1157.

7.Gokay S, Kendirci M, Kaynar L, Solmaz M, Cetin A, Kardas F, Soylu Ustkoyuncu P. Long-term efficacy of lipoprotein apheresis in the management of familial hypercholesterolaemia: Application of two different apheresis techniques in childhood. Transfus Apher Sci. 2016;54(2):282-288.

8.Ustkoyuncu PS, Ergül AB, Gökçek I, Hamsa H, Altuner Torun Y, Güven AS, Gökay S. A rare Lysosomal Storage Disease: Gaucher Type 2. JAMER 2017:2(3)13-20.

9.Ustkoyuncu PS, Kendirci M, Gökay S, Kardas F, Dursun I, Günes T. Effectiveness of Exchange Transfusion in Hyperlipoproteinemia Type 1. J Clin Anal Med 2017;8(suppl 3): 219-221.

10.Soylu Ustkoyuncu P, Mutlu FT, Kiraz A, Tag Balkis Z, Yel S. A Rare Cause of Neonatal Hemolytic Anemia: Glutathione Synthetase Deficiency. J Pediatr Hematol Oncol. 2018;40(1):45-49.

11. Ustkoyuncu PS, Güven AS, Kiraz A, Yılmaz A, Bozdemir ŞE, Gökay S. De Novo Mutation in ATP7A Gene with Severe Menkes Disease. Erciyes Med J. 2018; 40(2): 99-102

12.Ustkoyuncu PS, Kendirci M, Gökay S, Kardaş F.Hydroxy-3-methylglutaryl-CoA (HMG-CoA) Lyase Deficiency. Erciyes Med J. 2018; 40(3): 169-171.

13.Ustkoyuncu PS, Ergül AB, Altuner Torun Y. Investigation of Malnutrition in Hospitalized Children: A Point Prevalence Study from Kayseri. J Clin Anal Med 2018;9(1): 27-30.

14.Ustkoyuncu PS, Güven AS, Kiraz A, Gökay S, Doğan D. X-linked Adrenoleukodystrophy Initially Presenting with Severe Deafness. Turkiye Klinikleri J Case Rep. 2018;26(2):90-93.

15.Soylu Ustkoyuncu P, Nalcacioglu H, Bastug F, Yel S, Altuner Torun Y. Association of Mucopolysaccharidosis Type 4A and Bartter Syndrome. Iran J Kidney Dis. 2019;13(1):71-72.

16.Soylu Ustkoyuncu P, Kendirci M, Kardas F, Gokay S, Per H, Kacar Bayram A. Neutropenia and Increased Mean Corpuscular Volume (MCV) with Abnormal Neurologic Findings: A Case of Cobalamin D Deficiency. J Pediatr Hematol Oncol.2019;41(1):54-56.