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Education Information

Post Doctorate of Medicine, Hacettepe University, Tıp, Çocuk Sağlığı Ve Hastalıkları, Turkey 1996 - 2012
Expertise In Medicine, Hacettepe University, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı, Çocuk Metabolizma Hastalıkları, Turkey 1991 - 1996

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Child Health and Diseases

Academic Titles / Tasks

Professor, Hacettepe University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 1991 - Continues

Courses

Rikest, Undergraduate, 2017 - 2018
Fenilketonüri, Undergraduate, 2017 - 2018, 2016 - 2017
Metabolik hastalıklarda hipoglisemi, Undergraduate, 2017 - 2018, 2016 - 2017
Hypoglycemia in IEM, Undergraduate, 2017 - 2018, 2016 - 2017
Protein energy malnutrition, Undergraduate, 2017 - 2018, 2016 - 2017
Büyümenin değerlendirilmesi, Undergraduate, 2017 - 2018, 2016 - 2017
Phenylketonuria, Undergraduate, 2017 - 2018, 2016 - 2017
Problem based learning, Undergraduate, 2017 - 2018, 2016 - 2017
Probleme dayalı öğrenim, Undergraduate, 2017 - 2018, 2016 - 2017
Mukopolisakkaridoz, Undergraduate, 2017 - 2018
Mucopolysaccharidosis, Undergraduate, 2017 - 2018
Obesity and its prevention, Undergraduate, 2017 - 2018, 2016 - 2017
Obezite ve korunma yolları, Undergraduate, 2017 - 2018
Grotwh and its assessement, Undergraduate, 2017 - 2018
Rikets, Undergraduate, 2016 - 2017

Lizozomal depo hastalıkları : MPS örneği, Undergraduate, 2016 - 2017
Makronütrient eksiklikleri, Undergraduate, 2016 - 2017
Protein enerji malnütrisyonu, Undergraduate, 2016 - 2017
Obesite ve korunma yolları, Undergraduate, 2016 - 2017
Lysosomal storage disease: MPSs , Undergraduate, 2016 - 2017

Published journal articles indexed by SCI, SSCI, and AHCI

I. **High prevalence of low bone mineral density in young adults with phenylketonuria**
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IV. **Systematic literature review of the somatic comorbidities experienced by adults with phenylketonuria**
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VI. **The struggle that is phenylketonuria: What do the patients and caregivers suffer from**
DEMİREL D., SİVRİ S.
MEDICINE, no.25, 2024 (SCI-Expanded)

VII. **Predictors of eventual requirement of phenylalanine-restricted diet in young infants with phenylalanine hydroxylase deficiency initially managed with sapropterin monotherapy**
ÇIKI K., YILDIZ Y., KAHRAMAN A. B., ÖZGÜL R. K., COŞKUN T., DURSUN A., TOKATLI A., SİVRİ S.
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IX. **COVID-19 in inherited metabolic disorders: Clinical features and risk factors for disease severity**
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X. **Successful management of rhabdomyolysis with triheptanoin in a child with severe long-chain 3-hydroxyacyl-coenzyme A dehydrogenase (LCHAD) deficiency**
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XI. **Splenic Gaucheroma Leading to Incidental Diagnosis of Gaucher Disease in a 46-Year-Old Man with a**

Rare GBA Mutation: A Case Report
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XVI. **Does glutaric aciduria type 1 affect hearing function?**
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XVII. **Recommendations on phenylketonuria in Turkey**
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EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.63, no.11, 2020 (SCI-Expanded)

XXV. **Predictors of acute metabolic decompensation in children with maple syrup urine disease at the emergency department**
YILDIZ Y., Akcan Yildiz L., DURSUN A., TOKATLI A., COŞKUN T., TEKŞAM Ö., SİVRİ H. S.
EUROPEAN JOURNAL OF PEDIATRICS, vol.179, no.7, pp.1107-1114, 2020 (SCI-Expanded)

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YILDIZ Y., SİVRİ H. S.
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BALLIKAYA E., Eymirli P. S., YILDIZ Y., AVCU N., SİVRİ H. S., Uzamis-Tekcicek M.
TURKISH JOURNAL OF PEDIATRICS, vol.60, no.4, pp.400-406, 2018 (SCI-Expanded)

XXXVI. **Hyperinsulinemic Hypoglycemia in Congenital Disorder of Glycosylation Type-1a (CDG-1a)**
Vuralli D., YILDIZ Y., SİVRİ H. S., ALİKAŞİFOĞLU A.
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XXXVII. **Evaluation of spinal involvement in children with mucopolysaccharidosis VI: the role of MRI**
Bulut E., Pektas E., SİVRİ H. S., BİLGİNER B., UMAROĞLU M. M., Ozgen B.
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XXXVIII. **Efficacy, safety and population pharmacokinetics of sapropterin in PKU patients < 4 years: results from the SPARK open-label, multicentre, randomized phase IIIb trial**
Muntau A. C., Burlina A., Eyskens F., Freisinger P., De Laet C., Leuzzi V., Rutsch F., SİVRİ H. S., Vijay S., Bal M. O., et al.

ORPHANET JOURNAL OF RARE DISEASES, vol.12, 2017 (SCI-Expanded)

XXXIX. **Evaluation and identification of IDUA gene mutations in Turkish patients with mucopolysaccharidosis type I**
Atceken N., ÖZGÜL R. K., Yilmaz D. Y., TOKATLI A., COŞKUN T., SİVRİ H. S., DURSUN A., Karaca M.
TURKISH JOURNAL OF MEDICAL SCIENCES, vol.46, no.2, pp.404-408, 2016 (SCI-Expanded)

XL. **Late-diagnosed phenylketonuria in an eight-year-old boy with dyslexia and attention-deficit hyperactivity disorder**
YILDIZ Y., DURSUN A., TOKATLI A., COŞKUN T., SİVRİ H. S.
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ÖZLEM U., GÖKMEN ÖZEL H., COŞKUN T., ÖZGÜL R. K., YÜCEL YILMAZ D., BURCU H., TOKATLI A., DURSUN A., SİVRİ H. S.
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L. **Identification of Mutations and Evaluation of Cardiomyopathy in Turkish Patients with Primary Carnitine Deficiency**
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Supported Projects

SİVRİ S., Project Supported by Higher Education Institutions, 5. Uluslararası Pediatrik Hareket Bozuklulukları Sempozyumu, 2016 - 2017
SİVRİ H. S., Other International Funding Programs, SPARK EMR700773 003 BioMarin Study Number BMN 162 503, 2012 - 2017

Metrics

Publication: 121
Citation (WoS): 216
Citation (Scopus): 343
H-Index (WoS): 9
H-Index (Scopus): 12

Non Academic Experience

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