Asst. Prof. YILMAZ YILDIZ

Personal Information

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Education Information

Post Doctorate of Medicine, Hacettepe University, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, Turkey 2015 - 2018
Expertise In Medicine, Hacettepe Üniversitesi, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı, Turkey 2009 - 2014
Doctorate, Hacettepe University, Tıp Fakültesi (İngilizce), Temel Tıp Bilimleri Bölümü, Turkey 2007 - 2010

Foreign Languages

English, C2 Proficiency

Certificates, Courses and Trainings

Health&Medicine, Good Clinical Practice, çevrimiçi, 2018

Dissertations

Expertise In Medicine, Hiperfenilalaninemili hastaların gebelik sonuçlarının incelenmesi, Hacettepe University, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, 2014
Doctorate, Investigation of the role of Klf5 gene in muscle degeneration and differentiation, Hacettepe University, Sağlık Bilimleri Enstitüsü, Tıbbi Biyoloji A.B.D., 2010

Research Areas

Medicine, Health Sciences, Medical Biology, Internal Medicine Sciences, Child Health and Diseases, Pediatric Endocrinology and Metabolism

Academic Titles / Tasks

Assistant Professor, Hacettepe University, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, 2020 - Continues
Expert PhD, Pediatric Metabolic Diseases, 2018 - 2020
Research Assistant, Hacettepe University, Tıp Fakültesi (Türkçe), Dahili Tıp Bilimleri Bölümü, 2015 - 2018
Research Assistant, Hacettepe Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2009 - 2014

Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

I. **Recurrent Postinfectious Rhabdomyolysis in a 5-Year-Old Girl.**
Torun E., Yildiz Y., Yazici M., Çolak F., Kasapkara C.
Klinische Padiatrie, 2020 (Journal Indexed in SCI Expanded)

II. **A Rare Cause of Hyperinsulinemic Hypoglycemia: Kabuki Syndrome**
Mısırlıgil M., Yıldız Y., Akın O., Odabaşı G., Arslan M., Ünay B.
Journal of clinical research in pediatric endocrinology, 2020 (Journal Indexed in SCI Expanded)

III. **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, pp.1107-1114, 2020 (Journal Indexed in SCI)

IV. **Successful management of acute pancreatitis due to apolipoprotein C-II deficiency in a 37-day-old infant.**
Yıldız Y., Uysal Y., Çınar H., Özbay H., Kurt Ç., Kılıç M.
Pancreatology : official journal of the International Association of Pancreatology (IAP) ... [et al.], vol.20, pp.644-646, 2020 (Journal Indexed in SCI Expanded)

V. **Retrospective evaluation of 85 patients with urea cycle disorders: one center experience, three new mutations**
Nakip O. S. , YILDIZ Y., TOKATLI A.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.33, pp.721-728, 2020 (Journal Indexed in SCI)

VI. **Consensus guideline for the diagnosis and treatment of tetrahydrobiopterin (BH4) deficiencies**
Opladen T., Lopez-Laso E., Cortes-Saladelafont E., Pearson T. S. , SİVRİ H. S. , YILDIZ Y., Assmann B., Kurian M. A. , Leuzzi V., Heales S., et al.
ORPHANET JOURNAL OF RARE DISEASES, vol.15, 2020 (Journal Indexed in SCI)

VII. **Genotypes and estimated prevalence of phosphomannomutase 2 deficiency in Turkey differ significantly from those in Europe**
YILDIZ Y., Arslan M., Celik G., Kasapkara C. S. , Ceylaner S., DURSUN A., SİVRİ H. S. , COŞKUN T., TOKATLI A.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.182, pp.705-712, 2020 (Journal Indexed in SCI)

VIII. **Comment on: "Multiple acyl-CoA dehydrogenase deficiency in elderly carriers"**
YILDIZ Y., TOKATLI A.
JOURNAL OF NEUROLOGY, vol.267, pp.1209-1210, 2020 (Journal Indexed in SCI)

IX. **Oral health status of children with phenylketonuria**
BALLIKAYA E., YILDIZ Y., SİVRİ H. S. , TOKATLI A., DURSUN A., Olmez S., COŞKUN T., Tekcicek M. U.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.33, pp.361-365, 2020 (Journal Indexed in SCI)

X. **Two cases of vici syndrome presenting with corpus callosum agenesis, albinism, and severe developmental delay**
Hızal M., Yeke B., YILDIZ Y., Öztürk A., Gürbüz B. B. , COŞKUN T.
Turkish Journal of Pediatrics, vol.62, pp.474-478, 2020 (Journal Indexed in SCI Expanded)

XI. **Inborn errors of metabolism in the differential diagnosis of fatty liver disease**
YILDIZ Y., SİVRİ H. S.
TURKISH JOURNAL OF GASTROENTEROLOGY, vol.31, pp.3-16, 2020 (Journal Indexed in SCI)

XII. **When Pancreatitis Stinks A Rare Cause of Acute Recurrent Pancreatitis in a Child**
YILDIZ Y., Sahin G., Hosnut F. O.
PANCREAS, vol.49, 2020 (Journal Indexed in SCI)

XIII. **Rare cause of high anion gap metabolic acidosis in an infant: Succinyl-CoA:3-ketoacid transferase deficiency**
YILDIZ Y., Azapagasi E.
JOURNAL OF PAEDIATRICS AND CHILD HEALTH, vol.55, pp.1395-1396, 2019 (Journal Indexed in SCI)

XIV. **Determinants of Riboflavin Responsiveness in Multiple Acyl-CoA Dehydrogenase Deficiency**
YILDIZ Y., TALİM B., Haliloglu G., Topaloglu H., AKÇÖREN Z., DURSUN A., SİVRİ H. S. , COŞKUN T., TOKATLI A.

PEDIATRIC NEUROLOGY, vol.99, pp.69-75, 2019 (Journal Indexed in SCI)

XV. **Maternal phenylketonuria in Turkey: outcomes of 71 pregnancies and issues in management**
YILDIZ Y., SİVRİ H. S.
EUROPEAN JOURNAL OF PEDIATRICS, vol.178, pp.1005-1011, 2019 (Journal Indexed in SCI)

XVI. **Cognitive and behavioral impairment in mild hyperphenylalaninemia**
Evinç S., Pektaş E., Foto-Özdemir D., Yıldız Y., Karaboncuk Y., Bilginer-Gürbüz B., Dursun A., Tokatlı A., Coskun T., Öktem F., et al.
TURKISH JOURNAL OF PEDIATRICS, vol.60, pp.617-624, 2018 (Journal Indexed in SCI)

XVII. **Post-mortem detection of FLAD1 mutations in 2 Turkish siblings with hypotonia in early infancy**
YILDIZ Y., Olsen R. K. J. , SİVRİ H. S. , AKÇÖREN Z., Nygaard H. H. , TOKATLI A.
NEUROMUSCULAR DISORDERS, vol.28, pp.787-790, 2018 (Journal Indexed in SCI)

XVIII. **Oral health status in patients with mucopolysaccharidoses**
BALLIKAYA E., Eymirli P. S. , YILDIZ Y., AVCU N., SİVRİ H. S. , Uzamis-Tekcicek M.
TURKISH JOURNAL OF PEDIATRICS, vol.60, pp.400-406, 2018 (Journal Indexed in SCI)

XIX. **Hyperinsulinemic Hypoglycemia in Congenital Disorder of Glycosylation Type-1a (CDG-1a)**
Vuralli D., YILDIZ Y., SİVRİ H. S. , ALİKAŞİFOĞLU A.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.362-363, 2018 (Journal Indexed in SCI)

XX. **Hereditary Dopamine Transporter Deficiency Syndrome: Challenges in Diagnosis and Treatment**
YILDIZ Y., Pektas E., TOKATLI A., Haliloglu G.
NEUROPEDIATRICS, vol.48, pp.49-52, 2017 (Journal Indexed in SCI)

XXI. **Partial hydatidiform mole in a phenylketonuria patient treated with sapropterin dihydrochloride**
YILDIZ Y., DURSUN A., TOKATLI A., COŞKUN T., Sivri S.
GYNECOLOGICAL ENDOCRINOLOGY, vol.33, pp.19-20, 2017 (Journal Indexed in SCI)

XXII. **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.
TURKISH JOURNAL OF PEDIATRICS, vol.58, pp.94-96, 2016 (Journal Indexed in SCI)

XXIII. **Periostin is temporally expressed as an extracellular matrix component in skeletal muscle regeneration and differentiation**
OZDEMIR C., Akpulat U., SHARAFI P., YILDIZ Y., ONBAŞILAR İ., Kocaefe C.
Gene, vol.553, pp.130-139, 2014 (Journal Indexed in SCI Expanded)

XXIV. **Localized acute generalized exanthematous pustulosis with amoxicillin and clavulanic acid**
Ozkaya-Parlakay A., Azkur D., KARA A., YILDIZ Y., ORHAN D., CENGİZ A. B. , Ersoy-Evans S.
TURKISH JOURNAL OF PEDIATRICS, vol.53, pp.229-232, 2011 (Journal Indexed in SCI)

Articles Published in Other Journals

I. **Creatine Transporter Deficiency Presenting as Autism Spectrum Disorder**
Yıldız Y., Göçmen R., Yaramış A., Coşkun T., Haliloğlu G.
Pediatrics, vol.146, 2020 (Refereed Journals of Other Institutions)

II. **Infant Acute Lymphoblastic Leukemia with Atypical Presentation**
YAMAN BAJİN H. İ. , YILDIZ Y., akın ş., AYTAÇ EYÜPOĞLU Ş. S. , ÜNAL CANGÜL Ş., KUŞKONMAZ B. B. , Cetin M., SİVRİ H. S. , GÜMRÜK F.
Acta Medica, vol.50, pp.57-59, 2019 (Other Refereed National Journals)

III. **Mukopolisakkaridozlarda ortopedik sorunlar**
YILDIZ Y., SİVRİ H. S.
TOTBİD Dergisi, vol.15, pp.303-310, 2016 (Other Refereed National Journals)

IV. **Kalıtsal Metabolik Hastalıklarda Dental Bulgular**
YILDIZ Y., SİVRİ H. S.
Türkiye Klinikleri Çocuk Diş Hekimliği - Özel Konular, vol.2, pp.28-33, 2016 (Other Refereed National Journals)

Books & Book Chapters

I. **Nörotransmitter Bozukluklarında Klinik Yaklaşım**
Yıldız Y., Sivri H. S.
in: Kalıtsal Metabolik Hastalıklarda Hareket Bozuklukları, Dursun,Ali, Editor, Türkiye Klinikleri Yayınevi, Ankara, pp.20-29, 2020

Supported Projects

Auricchio A., Sivri H. S. , Yıldız Y., FP7 Project, MEUSix Ağır bir lizozomal depo hastalığı olan MPS tip VI lı hastalarda gen tedavisi, 2014 - 2017

Scientific Refereeing

TURKISH JOURNAL OF PEDIATRICS, Journal Indexed in SCI-E, August 2020
JOURNAL OF INTEGRATIVE NEUROSCIENCE, Journal Indexed in SCI-E, June 2020
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, Journal Indexed in SCI-E, April 2020
ANNALS OF INDIAN ACADEMY OF NEUROLOGY, Journal Indexed in SCI-E, August 2019
EUROPEAN JOURNAL OF PEDIATRICS, Journal Indexed in SCI-E, June 2019
Project Supported by Private Organizations in Other Countries, Telethon Foundation, Italy, June 2019

Citations

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