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Published journal articles indexed by SCI, SSCI, and AHCI

- I. **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.
Molecular Genetics and Metabolism, vol.140, no.3, 2023 (SCI-Expanded)
- II. **COVID-19 in inherited metabolic disorders: Clinical features and risk factors for disease severity**
KAHRAMAN A. B., YILDIZ Y., ÇIKI K., Erdal I., AKAR H. T., DURSUN A., TOKATLI A., SİVRİ S.
Molecular Genetics and Metabolism, vol.139, no.2, 2023 (SCI-Expanded)
- III. **Efficacy of Tele-CO-OP in Children With Organic Acidemia: A Pilot Randomized Controlled Trial**
Dursun E. L., BUMİN G., ÇIKI K., SİVRİ S.
OTJR Occupation, Participation and Health, 2023 (SSCI)
- IV. **Vascular and structural analyses of retinal and choroidal alterations in Fabry disease: the effect of hyperreflective foci and retinal vascular tortuosity**
YANIK ODABAŞ Ö., ÇIKI K., ÖZMERT E., Sivri S.
OPHTHALMIC GENETICS, vol.43, no.3, pp.344-353, 2022 (SCI-Expanded)
- V. **Evaluation of sleep-disordered breathing and its relationship with respiratory parameters in children with mucopolysaccharidosis Type IVA and VI**
Ademhan Tural D., EMİRALİOĞLU N., DOĞRU ERSÖZ D., Ozsezen B., Ipek O. F., SUNMAN B., Nayir Buyuksahin H., GÜZELKAŞ İ., ÇIKI K., Kilic K., et al.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.185, no.8, pp.2306-2314, 2021 (SCI-Expanded)
- VI. **Caring for a Child with Phenylketonuria: Parental Experiences from a Eurasian Country**
ZENGİN AKKUŞ P., BİLGİNER GÜRBÜZ B., ÇIKI K., İLTER BAHADUR E., KARAHAN S., ÖZMERT E. N., COŞKUN T., Sivri S.
JOURNAL OF DEVELOPMENTAL AND BEHAVIORAL PEDIATRICS, vol.41, no.3, pp.195-202, 2020 (SCI-Expanded)
- VII. **A rare cause of acute abdominal pain in a patient with Primary ciliary dyskinesia with situs inversus totalis**
Ciki K., Turer Ö., Hizal M., Tugcu G. D., EMİRALİOĞLU N., Yalcin E., Ersoz D., Kiper N., Ozcelik U.
TURKISH JOURNAL OF PEDIATRICS, vol.62, no.1, pp.156-159, 2020 (SCI-Expanded)
- VIII. **Pulmonary complications following hematopoietic stem cell transplantation in children**
ÇIKI K., DOĞRU ERSÖZ D., Kuskonmaz B. B., EMİRALİOĞLU N., Yalcin E., Ozcelik U., Uckan-Cetinkaya D., Kiper N.
TURKISH JOURNAL OF PEDIATRICS, vol.61, no.1, pp.59-70, 2019 (SCI-Expanded)
- IX. **A fatal Clostridium perfringens infection with hemolysis after chemotherapy in an adolescent**
Kukul M. G., Ciki K., Karadag-Oncel E., CENGİZ A. B., Kuskonmaz B. B., Tavail B., KESİCİ S.
ARCHIVOS ARGENTINOS DE PEDIATRIA, vol.115, no.2, 2017 (SCI-Expanded)

Articles Published in Other Journals

- I. **The Complication of Endotracheal Intubation in a Patient with Mucopolysaccharidosis Type IIIA**

Tural D. A., EMİRALİOĞLU N., Ozsezen B., ÇIKI K., Ozcan N., ARDIÇLI B., Sivri S.

INTERNATIONAL JOURNAL OF PEDIATRICS-MASHHAD, vol.9, no.2, pp.12909-12913, 2021 (ESCI)

II. Cardiomyopathy in patients with type 1 tyrosinemia, and the effect of nitisinone treatment on cardiomyopathy

BİLGİNER GÜRBÜZ B., AYKAN H. H., ÇIKI K., KARAGÖZ T., Sivri S., DURSUN A., TOKATLI A., COŞKUN T.

CUKUROVA MEDICAL JOURNAL, vol.46, no.4, pp.1419-1425, 2021 (ESCI)

III. Clinical evaluation of patients with congenital lung malformations Konjenital akciğer malformasyonu olan hastaların klinik değerlendirilmesi

EMİRALİOĞLU N., Çiki K., Tugcu G. D., Yalçın E., Ersöz D., Kiper N., Özçelik U.

Cocuk Sagligi ve Hastaliklari Dergisi, vol.60, no.1, pp.1-6, 2017 (Scopus)

Refereed Congress / Symposium Publications in Proceedings

I. Listening Parents Caring A Child With Phenylketonuria

ZENGİN AKKUŞ P., BİLGİNER GÜRBÜZ B., ÇIKI K., İLTER BAHADUR E., KARAHAN S., ÖZMERT E. N., COŞKUN T., SİVRİ H. S.

3rd International Developmental Pediatrics Association Congress, 9 - 12 December 2019

II. Fenilketonüri ile Yaşamak: Anne Ve Babaların Penceresinden

ZENGİN AKKUŞ P., BİLGİNER GÜRBÜZ B., ÇIKI K., İLTER BAHADUR E., KARAHAN S., ÖZMERT E. N., COŞKUN T., SİVRİ H. S.

63. Türkiye Milli Pediatri Kongresi, Turkey, 30 October - 03 November 2019

III. Hyperphenylalaninemia due to novel JCDNA12 mutation

SİVRİ H. S., ÇIKI K., YÜCEL YILMAZ D., GÜRSES CİLA H. E., ÖZGÜL R. K., TOKATLI A., COŞKUN T., DURSUN A.

SSIEM 2019: Annual Symposium of the Society for the Study of Inborn Errors of Metabolism, Rotterdam, Netherlands, 3 - 06 September 2019, vol.42, pp.324

IV. Antibiotic resistance of Haemophilus influenzae in primary Ciliary Dyskinesia patients

ÇIKI K., SANCAK B., DEMİRCİ S., EMİRALİOĞLU N., TUĞCU G. D., EŞREF S., HIZAL M., YALÇIN E., DOĞRU ERSÖZ D., KİPER E. N., et al.

2nd BEAT-PCD Conference and 3rd PCD training School, Valencia, Spain, 18 - 21 April 2017

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