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Education Information

Post Doctorate of Medicine, Cukurova University, Tıp Fakültesi, Çocuk Endokrinoloji Ve Metabolizma Bilim Dalı, Turkey 1995 - 2005
Expertise In Medicine, Cukurova University, Tıp Fakültesi, Pediatri Anabilim Dalı, Turkey 1990 - 1995
Undergraduate, Cukurova University, Tıp Akültesi, Turkey 1982 - 1988

Foreign Languages

English, B2 Upper Intermediate

Research Areas

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

Academic Titles / Tasks

Professor, Cukurova University, Tıp Fakültesi, Dahili Tıp Bilimleri, 1996 - Continues

Academic and Administrative Experience

Cukurova University, Hemşirelik Myo, 2001 - 2002

Courses

Metabolik karaciğer hastalıkları , Undergraduate, 2016 - 2017

Advising Theses

ÖNENLİ MUNGAN H., Primer ailesel hiperlipidemi tanılı çocuk ve adölesanlarda demografik, klinik özellikler ve bunların mutasyon analizi sonuçlarıyla ilişkisi, Expertise In Medicine, i.kaplan(Student), 2016
ÖNENLİ MUNGAN H., Çukurova Üniversitesi Tıp Fakültesi Pediatrik Metabolizma Hastalıkları ve Beslenme Polikliniğinde tanı alan veya takibe giren kalıtsal metabolik hastalığı olan hastaların tanılarının, klinik ve laboratuar bulgularının analizi ile takip sonuçlarının değerlendirilmesi, Expertise In Medicine, a.kara(Student), 2012
ÖNENLİ MUNGAN H., Çocukluk çağı obezitesinde metabolik parametrelerin diyet ve egzersizle ilişkisi, Expertise In Medicine, i.öncü(Student), 2009

Jury Memberships

Associate Professor Exam, Doçentlik sınavı, Doçentlik sınavı, June, 2016

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Real-world patient data on immunity and COVID-19 status of patients with MPS, Gaucher, and Pompe diseases from Turkey**
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IV. **Investigating myelin oligodendrocyte glycoprotein antibodies in hereditary citrullinemia**
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V. **Dysarthria, Ataxia, and Dystonia Associated with COX20 (FAM36A) Gene Mutation: A Case Report of a Turkish Child**
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VI. **Defining clinical subgroups and genotype-phenotype correlations in NBAS-associated disease across 110 patients**
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VIII. **A possible biomarker of neurocytolysis in infantile gangliosidoses: aspartate transaminase**
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XIII. **A Case Report of a Very Rare Association of Tyrosinemia type I and Pancreatitis Mimicking Neurologic Crisis of Tyrosinemia Type I.**
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XVI. **Brown-Vialetto-Van Laere syndrome: two siblings with a new mutation and dramatic therapeutic effect of high-dose riboflavin.**
HOROZ O., Mungan N., YILDIZDAS D., HERGÜNER Ö., CEYLANER S., KOR D., WATERHAM H., COSKUN T.
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XVIII. **A Desensitization Method to Maintain Enzyme Replacement Therapy in Mucopolysaccharidosis Type VI.**
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XIX. **Genotypic and phenotypic features of the cystinosis patients from the South Eastern part of Turkey.**
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XX. **Primary systemic carnitine deficiency: a Turkish case with a novel homozygous SLC22A5 mutation and 14 years follow-up.**
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XXII. **A 17-Year-Old Girl with Chronic Intermittent Abdominal Pain**
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PEDIATRIC ANNALS, vol.44, no.4, pp.139-141, 2015 (SCI-Expanded)

XXIII. **TYROSINEMIA TYPE 1 AND NEUROGENIC CRISIS: A CASE REPORT**
BULUT F. D., KOR D., Onenli-Mungan N., Yükselmiş U., İNCECİK F., Yildizdas D.
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GÜRBÜZ F., KOTAN L. D., Mengen E., ŞIKLAR Z., BERBEROĞLU M., Dokmetas S., Kilicli M. F., Guven A., KİREL B., Saka N., et al.
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XXV. **Serum IGF-1 and IGFBP-3 Levels in Healthy Children Between 0 and 6 Years of Age**
YÜKSEL B., Ozbek M. N., Mungan N. O., Darendeliler F., Budan B., BİDECİ A., Cetinkaya E., BERBEROĞLU M., Evliyaoglu O., Yesilkaya E., et al.
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XXVI. **Ambulatory Blood Pressure Monitoring and Serum Nitric Oxide Concentration in Type 1 Diabetic Children**
Horoz O. O., YÜKSEL B., Bayazit A., ATTILA G., SERTDEMİR Y., Mungan N. O., TOPALOĞLU A. K., OZER G.
ENDOCRINE JOURNAL, vol.56, no.3, pp.477-485, 2009 (SCI-Expanded)

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YÜKSEL B., Ozbek M. N., Darendeliler F., BİDECİ A., Cetinkaya E., BERBEROĞLU M., Evliyaoglu O., Bas F., Mungan N. O., Yesilkaya E., et al.
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XXVIII. **A homozygous recurring mutation in WISP3 causing progressive pseudorheumatoid arthropathy of childhood**
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İNCECİK F., Herguner O. M., Mungan N. O.
JOURNAL OF PEDIATRIC NEUROSCIENCES, vol.15, no.2, pp.86-89, 2020 (ESCI)

II. **Vitamin B12 levels in patients with mucopolysaccharidosis**
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CUKUROVA MEDICAL JOURNAL, vol.45, no.2, pp.401-407, 2020 (ESCI)

III. **A 6-Month-Old Boy with Reddish, Scaly Skin: Netherton Syndrome**
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IV. **Neurological assessment of 38 late-diagnosed children with classic phenylketonuria**
HAYTOĞLU Z., Herguner O., SOYUPAK S., TOPALOĞLU A. K., YÜKSEL B., OZER G., Mungan H. N. O.
CUKUROVA MEDICAL JOURNAL, vol.41, no.1, pp.21-27, 2016 (ESCI)

V. **The analysis of the phenylalanine hydroxylase gene mutations by sequencing and ARMS techniques**

in Turkish patients
Luleyap U., PAZARCI P., CÖMERTPAY G., Onenli H. N., PAZARBAŞI A., ALPTEKİN D., KASAP H., FROSTER U.
CUKUROVA MEDICAL JOURNAL, vol.41, no.4, pp.702-708, 2016 (ESCI)

VI. **Brown-Vialetto-Van Laere syndrome: Two siblings with a new mutation and dramatic therapeutic effect of high dose riboflavin**
ÖNENLİ MUNGAN H.
J PEDIATR ENDOCRINOL METAB, vol.2015, pp.198-205, 2015 (Peer-Reviewed Journal)

VII. **Continuous Venovenous Hemodiafiltration in Three Newborn Patients with Hyperammonemia**
YAPICIOĞLU YILDIZDAŞ H., Yildizdas D., GÜLDEREN ÖZLÜ F., Mert K., Mungan N. O.
CUKUROVA MEDICAL JOURNAL, vol.40, pp.161-166, 2015 (ESCI)

VIII. **Fabry Disease: A Turkish Case with a Novel Mutation and Dermatological Manifestations**
Mungan N. O., Temiz F., Yilmaz B. S., Ozbek M. N., KARAKAŞ M., TOPALOĞLU A. K., YÜKSEL B.
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IX. **Evaluation of Two Different Pamidronate Treatment Protocols in Children with Osteogenesis Imperfecta**
ÖNENLİ MUNGAN H., gürbüz F., mengen e., özgür ö., topaloğlu a. k., yüksel b.
CUKUROVA MEDICAL JOURNAL, vol.39, no.3, pp.532-539, 2014 (Peer-Reviewed Journal)

X. **Hashimoto's Encephalopathy: Four Cases and Review of Literature.**
YÜKSEL B., İNCECİK F., HERGÜNER M., ÖZGÜR HOROZ Ö., YILDIZDAŞ R. D., ALTUNBAŞAK Ş., ÖNENLİ MUNGAN H., GÜL MERT G.
Int J Neurosci, vol.0, 2013 (Peer-Reviewed Journal)

XI. **Autism symptoms related to Tyrosinemia type III: a case report**
YOLGA TAHİROĞLU A., ÖNENLİ MUNGAN H., FIRAT S., AVCI A.
TURKISH JOURNAL OF ENDOCRINOLOGY AND METABOLISM, vol.12, no.2, pp.55-56, 2008 (ESCI)

XII. **Pediatrik kafa travmalarında idrar antidiürik hormon seviyeleri**
YAMAN A., YÜKSEL B., AKSARAY N., ÖNENLİ MUNGAN H., YILDIZDAŞ R. D., Alhan A.
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Refereed Congress / Symposium Publications in Proceedings

I. **YENİDOĞANDA BİYOKİMYASAL HİPERTROİDİSM**
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IV. **First case report of Gaucher disease and Graves' thyroiditis**
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V. **SON DÖNEM BÖBREK YETMEZLİĞİ VE SİSTİNOZİS: 41 YAŞINDA GÖZ MUAYENESİ İLE TANI ALABİLEN BİR OLSU SUNUMU**
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VII. **Ağır laktik asidoz gelişen MMA?lı bir olguda yüksek doz askorbik asit tedavisiyle salah**
KÖR D., ÖNENLİ MUNGAN H., Özgür Ö., Yükselmiş U., Sarı Y., Ece Ü., Güntek S., YILDIZDAŞ R. D.
. XII. Metabolik Hastalıklar ve Beslenme Kongresi., Eskişehir, Turkey, 1 - 04 May 2013

VIII. **Ağır laktik asidoz gelişen MMA?lı bir olguda yüksek doz askorbik asit tedavisiyle salah**
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IX. **Brown-Vialetto-Van Laere ve Fazio Londe Sendromlu Bir Olgu Sunumu.**
Gülen M., YILDIZDAŞ R. D., Sarı M., ÖZGÜR HOROZ Ö., Yükselmiş U., ÖNENLİ MUNGAN H.
Uluslararası Katılımlı X. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Kongresi VI. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Hemşireliği Kongresi, Antalya, Turkey, 3 - 07 April 2013

X. **Brown-Vialetto-Van Laere ve Fazio Londe Sendromlu Bir Olgu Sunumu.**
Gülen M., YILDIZDAŞ R. D., Sarı M., ÖZGÜR HOROZ Ö., Yükselmiş U., ÖNENLİ MUNGAN H.
Uluslararası Katılımlı X. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Kongresi VI. Ulusal Çocuk Acil Tıp ve Yoğun Bakım Hemşireliği Kongresi, Antalya, Turkey, 3 - 07 April 2013

XI. **"A New Method For The Detection Of Ivs 10 11g-A Mutation In Phenylalanine Hydroxilase Gene With Arms Technique In Turkish Phenylketonuria Patients"**
LÜLEYAP H. Ü., ONATOGLU D., PAZARBAŞI A., GUZEL A. I., OZER G., ÖNENLİ MUNGAN H., KASAP M., ALPTEKİN D.
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Supported Projects

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Metrics

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