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Personal Information

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Education Information

Post Doctorate of Medicine, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 2005 - 2009

Expertise In Medicine, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 1999 - 2004

Under Graduate, Istanbul University, İstanbul Tıp Fakültesi, İstanbul Tıp Pr., Turkey 1992 - 1998

Academic Titles / Tasks

Associate Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2014 - Continues

Expert, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2014 - 2014

Lecturer, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2009 - 2011

Professional Experience

Vice Dean, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2020 - Continues

Fakülte Kurulu Üyesi, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2018 - Continues

Fakülte Yönetim Kurulu Üyesi, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2018 - Continues

Ethics Committee Member, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2019 - 2020

Yıl Koordinatörü, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri, 2015 - 2020

Courses

VİTAMİN VE MİNERAL BOZUKLUKLARI, Under Graduate, 2016 - 2017, 2017 - 2018

NE ZAMAN METABOLİK HASTALIK DÜŞÜNELİM?, Under Graduate, 2016 - 2017, 2017 - 2018

VİTAMİN VE MİNERAL BOZUKLUKLARI, Doctorate, 2017 - 2018

BESLENMEYE GİRİŞ, Under Graduate, 2016 - 2017, 2017 - 2018

Advising Theses

OKUR İ., İdiyopatik proteinürisi olan çocuk hastalarda alfa-galaktosidaz a eksikliğinin (fabry hastalığının) taranması, Expertise In Medicine, M.AKGÜL(Student), 2018

OKUR İ., 2000-2013 yılları arasında Gazi Üniversitesi Hastanesinde yatan ve total pranteral nutrisyon uygulanan çocuk hastaların değerlendirilmesi, Expertise In Medicine, B.TANRIKULU(Student), 2016

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

1. Nutritional Status of Syrian Refugees in Early Adolescence Living in Turkey

PEHLİVANTÜRK KIZILKAN M., ÖZMERT E. N. , DERMAN O., OKUR İ., KAYNAK M. O. , Adiguzel A., Sahan-Seref I., Kurekci F., BİDECİ A., Hasanoglu E.

JOURNAL OF IMMIGRANT AND MINORITY HEALTH, vol.22, pp.1149-1154, 2020 (Journal Indexed in SSCI)

- II. **Beneficial Effects of Modified Atkins Diet in Glycogen Storage Disease Type IIIa**
Olgac A., İNCİ A., OKUR İ., BİBEROĞLU G., Oguz D., EZGÜ F. S. , Kasapkara C. S. , Aktas E., TÜMER L.
ANNALS OF NUTRITION AND METABOLISM, vol.76, pp.233-241, 2020 (Journal Indexed in SCI)
- III. **CDH13andLPHN3Gene Polymorphisms in Attention-Deficit/Hyperactivity Disorder: Their Relation to Clinical Characteristics**
ÖZASLAN A., GÜNEY E., ERGÜN M. A. , OKUR İ., YAPAR D.
JOURNAL OF MOLECULAR NEUROSCIENCE, 2020 (Journal Indexed in SCI)
- IV. **Vitamin D Levels and Bone Mineral Density in Inborn Errors of Metabolism Requiring Specialised Diets**
Olgac A., İNCİ A., OKUR İ., Ezgu F., BİBEROĞLU G., Turner L.
JCPSP-JOURNAL OF THE COLLEGE OF PHYSICIANS AND SURGEONS PAKISTAN, vol.29, pp.1207-1211, 2019 (Journal Indexed in SCI)
- V. **High incidence of co-existing factors significantly modifying the phenotype in patients with Fabry disease**
Koca S., TÜMER L., OKUR İ., ERTEN Y., Bakkaloglu S., BİBEROĞLU G., Kasapkara C., Kucukcongar A., DALGIÇ B., ÖZHAN OKTAR S., et al.
GENE, vol.687, pp.280-288, 2019 (Journal Indexed in SCI)
- VI. **Epilepsy in Biotinidase Deficiency Is Distinct from Early Myoclonic Encephalopathy**
Guliyeva U., OKUR İ., Dulac O., Khalilov O., Guliyeva S.
NEUROPEDIATRICS, vol.49, pp.417-418, 2018 (Journal Indexed in SCI)
- VII. **Hematologic Findings of Inherited Metabolic Disease: They are More Than Expected**
Sal E., Yenicesu I., OKUR İ., KAYA Z., EZGÜ F. S. , KOÇAK Ü., TÜMER L., Gursel T., Hasanoglu A.
JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.40, pp.355-359, 2018 (Journal Indexed in SCI)
- VIII. **Patient With Niemann-Pick Type C Presenting With a Jaw Mass Characterized With Lymph Node Involvement by Niemann-Pick Cells**
İNCİ A., OKUR İ., ESENDAĞLI G., OKUR A., Olgac A., EZGÜ F. S. , TÜMER L.
JOURNAL OF PEDIATRIC HEMATOLOGY ONCOLOGY, vol.40, pp.243-245, 2018 (Journal Indexed in SCI)
- IX. **Structure-function analyses of microsomal triglyceride transfer protein missense mutations in abetalipoproteinemia and hypobetalipoproteinemia subjects**
Walsh M. T. , Di Leo E., OKUR İ., Tarugi P., Hussain M. M.
BIOCHIMICA ET BIOPHYSICA ACTA-MOLECULAR AND CELL BIOLOGY OF LIPIDS, vol.1861, pp.1623-1633, 2016 (Journal Indexed in SCI)
- X. **Bi-allelic Mutations in KLHL7 Cause a Crisponi/CISS1-like Phenotype Associated with Early-Onset Retinitis Pigmentosa**
Angius A., Uva P., Buers I., Oppo M., Puddu A., Onano S., Persico I., Loi A., Marcia L., Hoehne W., et al.
AMERICAN JOURNAL OF HUMAN GENETICS, vol.99, pp.236-245, 2016 (Journal Indexed in SCI)
- XI. **Audiologic evaluations of children with mucopolysaccharidosis**
Gokdogan C., ALTINYAY Ş., Gokdogan O., TUTAR H., GÜNDÜZ B., OKUR İ., TÜMER L., KEMALOĞLU Y. K.
BRAZILIAN JOURNAL OF OTORHINOLARYNGOLOGY, vol.82, pp.281-284, 2016 (Journal Indexed in SCI)
- XII. **The Janus-faced manifestations of homozygous familial hypobetalipoproteinemia due to apolipoprotein B truncations**
Di Leo E., Eminoglu T., Magnolo L., Bolkent M. G. , TÜMER L., OKUR İ., Tarugi P.
JOURNAL OF CLINICAL LIPIDOLOGY, vol.9, pp.400-405, 2015 (Journal Indexed in SCI)
- XIII. **Monocarboxylate Transporter 1 Deficiency and Ketone Utilization**
van Hasselt P. M. , Ferdinandusse S., Monroe G. R. , Ruiten J. P. N. , Turkenburg M., Geerlings M. J. , Duran K., Harakalova M., van der Zwaag B., Monavari A. A. , et al.
NEW ENGLAND JOURNAL OF MEDICINE, vol.371, pp.1900-1907, 2014 (Journal Indexed in SCI)
- XIV. **Serum dipeptidyl peptidase-IV: A better screening test for early detection of**

mucopolysaccharidosis?

Kurt I, Sertoglu E, OKUR İ, Tapan S, Uyanik M, Kayadibi H, EZGÜ F. S. , Aydin H. I. , Hasanoglu A.
CLINICA CHIMICA ACTA, vol.431, pp.250-254, 2014 (Journal Indexed in SCI)

- XV. **Screening for Fabry disease in patients undergoing dialysis for chronic renal failure in Turkey: Identification of new case with novel mutation**
OKUR İ, Ezgu F, BİBEROĞLU G, Turner L, ERTEN Y, Isitman M, Eminoglu F. T. , Hasanoglu A.
GENE, vol.527, pp.42-47, 2013 (Journal Indexed in SCI)
- XVI. **Quality of life in children treated with restrictive diet for inherited metabolic disease**
Eminoglu T. F. , Soysal S. A. , TÜMER L., OKUR İ, Hasanoglu A.
PEDIATRICS INTERNATIONAL, vol.55, pp.428-433, 2013 (Journal Indexed in SCI)
- XVII. **Oxidized low-density lipoprotein levels and carotid intima-media thickness as markers of early atherosclerosis in prepubertal obese children**
OKUR İ, TÜMER L., EZGÜ F. S. , Yesilkaya E, Aral A., ÖZHAN OKTAR S., BİDECİ A., Hasanoglu A.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.26, pp.657-662, 2013 (Journal Indexed in SCI)
- XVIII. **Hypercalcemia in glycogen storage disease type I patients of Turkish origin**
Kasapkara C. S. , TÜMER L., OKUR İ, Eminoglu T, EZGÜ F. S. , Hasanoglu A.
TURKISH JOURNAL OF PEDIATRICS, vol.54, pp.35-37, 2012 (Journal Indexed in SCI)
- XIX. **The levels of asymmetric dimethylarginine, homocysteine and carotid intima-media thickness in hypercholesterolemic children**
Hasanoglu A, OKUR İ, Oren A. C. , BİBEROĞLU G., ÖZHAN OKTAR S., Eminoglu F. T. , TÜMER L.
TURKISH JOURNAL OF PEDIATRICS, vol.53, pp.522-527, 2011 (Journal Indexed in SCI)
- XX. **N-carbamylglutamate treatment for acute neonatal hyperammonemia in isovaleric acidemia**
Kasapkara C. S. , EZGÜ F. S. , OKUR İ, TÜMER L., BİBEROĞLU G., Hasanoglu A.
EUROPEAN JOURNAL OF PEDIATRICS, vol.170, pp.799-801, 2011 (Journal Indexed in SCI)
- XXI. **IDUA Mutational Profiling of a Cohort of 102 European Patients with Mucopolysaccharidosis Type I: Identification and Characterization of 35 Novel alpha-L-iduronidase (IDUA) Alleles**
Bertola F., Filocamo M., Casati G., Mort M., Rosano C., Tytki-Szymanska A., Tuysuz B., Gabrielli O., Grossi S., Scarpa M., et al.
HUMAN MUTATION, vol.32, 2011 (Journal Indexed in SCI)
- XXII. **Harderoporphyria due to homozygosity for coproporphyrinogen oxidase missense mutation H327R**
Hasanoglu A, Balwani M, Kasapkara C. S. , EZGÜ F. S. , OKUR İ, TÜMER L., Cakmak A., Nazarenko I, Yu C., Clavero S., et al.
JOURNAL OF INHERITED METABOLIC DISEASE, vol.34, pp.225-231, 2011 (Journal Indexed in SCI)
- XXIII. **AN INFANTILE CASE OF ZELLWEGER SYNDROME PRESENTED WITH KABUKI-LIKE PHENOTYPE**
Ezgu F, Eminoglu T, OKUR İ, Gunduz M, Tumer L, Hasanoglu A, Dalgic B.
GENETIC COUNSELING, vol.22, pp.217-220, 2011 (Journal Indexed in SCI)
- XXIV. **The role of viral agents in aetiopathogenesis of acute rheumatic fever**
Olgunturk R, OKUR İ, Cirak M. Y. , OĞUZ A. D. , Akalin N, Turet S, Tunaoglu S.
CLINICAL RHEUMATOLOGY, vol.30, pp.15-20, 2011 (Journal Indexed in SCI)
- XXV. **A NOVEL MUTATION OF THE CLAUDIN 16 GENE IN FAMILIAL HYPOMAGNESEMIA WITH HYPERCALCIURIA AND NEPHROCALCINOSIS MIMICKING RICKETS**
Kasapkara C. S. , Tumer L., OKUR İ, Hasanoglu A.
GENETIC COUNSELING, vol.22, pp.187-192, 2011 (Journal Indexed in SCI)
- XXVI. **3-Methylcrotonyl-CoA Carboxylase Deficiency: Phenotypic Variability in a Family**
Eminoglu F. T. , Ozcelik A. A. , OKUR İ, TÜMER L., BİBEROĞLU G., DEMİR E., Hasanoglu A., Baumgartner M. R.
JOURNAL OF CHILD NEUROLOGY, vol.24, pp.478-481, 2009 (Journal Indexed in SCI)
- XXVII. **Crisponi Syndrome: A New Case With Additional Features and New Mutation in CRLF1**
OKUR İ, TÜMER L., Crisponi L., Eminoglu F. T. , Chiappe F., CİNAZ P., Yenicesu I, Hasanoglu A.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, pp.3237-3239, 2008 (Journal Indexed in SCI)
- XXVIII. **Multisystem involvement in a patient due to accumulation of amylopectin-like material with diminished branching enzyme activity**

- Eminoglu T. F. , Tumer L., OKUR İ., Olgunturk R., Hasanoglu A., Gonul I. I. , Dalgic B.
JOURNAL OF INHERITED METABOLIC DISEASE, vol.31, 2008 (Journal Indexed in SCI)
- XXIX. **Lipid apheresis applications in childhood: Experience in the University Hospital of Gazi**
Eminoglu T. F. , Yenicesu I., TÜMER L., OKUR İ., Dilsiz G., Hasanoglu A.
TRANSFUSION AND APHERESIS SCIENCE, vol.39, pp.235-240, 2008 (Journal Indexed in SCI)
- XXX. **Rapid screening of 10 common mutations in Turkish Gaucher patients using electronic DNA microarray**
Ezgu F., Hasanoglu A., OKUR İ., Biberoglu G., Tumer L., Eminoglu T., Dogan H.
BLOOD CELLS MOLECULES AND DISEASES, vol.40, pp.246-247, 2008 (Journal Indexed in SCI)
- XXXI. **Vitamin D intoxication and hypercalcaemia in an infant treated with pamidronate infusions**
Ezgu F., Buyan N., Gunduz M., Tumer L., OKUR İ., Hasanoglu A.
EUROPEAN JOURNAL OF PEDIATRICS, vol.163, pp.163-165, 2004 (Journal Indexed in SCI)

Articles Published in Other Journals

- I. **Citrullinemia with an Atypical Presentation: Paroxysmal Hypoventilation Attacks**
Ozturk Z., HIRFANOĞLU T., İNCİ A., OKUR İ., KOÇ E., TÜMER L., ARHAN E., Aydın K., SERDAROĞLU A.
JOURNAL OF PEDIATRIC NEUROSCIENCES, vol.13, pp.276-278, 2018 (Journal Indexed in ESCI)
- II. **Association Between Soluble CD40 Ligand and Hypercholesterolemia in Children and Adolescents**
Yavas A. K. , Eminoglu T. F. , OKUR İ., Aral A., Hasanoglu A., TÜMER L.
JOURNAL OF PEDIATRIC RESEARCH, vol.4, pp.1-5, 2017 (Journal Indexed in ESCI)
- III. **Giant bronchogenic cyst mimicking tension pneumothorax.**
Demircan S., Kurul İ. C. , Tokur M., Memis L., Okur İ.
Asian cardiovascular & thoracic annals, vol.14, pp.244-6, 2006 (Refereed Journals of Other Institutions)
- IV. **Karbonmonoksit Zehirlenmesinde Hiperbarik Oksijen Tedavisi İki Olgu Sunumu**
OKUR İ., SERDAROĞLU A., OKUR A., BUYAN N., DÜNDAR K., ARGAN M., ÖZDEMİR B., GÜCÜYENER K.
TÜRKİYE KLİNİKLERİ J PEDIATR, vol.14, pp.220-222, 2005 (Other Refereed National Journals)

Books & Book Chapters

- I. **Glikojen Depo Hastalıkları**
HASANOĞLU A., OKUR İ.
in: TEMEL PEDIATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ TIP KİTAPEVLERİ, Ankara, pp.1107-1111, 2020
- II. **Tirozin Metabolizması Bozuklukları**
OKUR İ.
in: TEMEL PEDIATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ TIP KİTAPEVLERİ, Ankara, pp.1144-1148, 2020
- III. **Keton Cisim Yapım ve Yıkım Bozuklukları**
OKUR İ.
in: TEMEL PEDIATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ TIP KİTAPEVLERİ, Ankara, pp.1132-1135, 2020
- IV. **Yağ Asidi Oksidasyon Bozuklukları**
İNCİ A., OKUR İ.
in: TEMEL PEDIATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ TIP KİTAPEVLERİ, Ankara, pp.1126-1130, 2020
- V. **Mitokondriyal Hastalıklar**
OKUR İ.
in: TEMEL PEDIATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ

TIP KİTABEVLERİ, Ankara, pp.1136-1139, 2020

- VI. **Fruktoz Metabolizması Bozuklukları**
HASANOĞLU A., OKUR İ.
in: TEMEL PEDİATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ
TIP KİTABEVLERİ, Ankara, pp.1122-1123, 2020
- VII. **Metionin Metabolizması Bozuklukları**
OKUR İ.
in: TEMEL PEDİATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ
TIP KİTABEVLERİ, Ankara, pp.1156-1158, 2020
- VIII. **Vitaminler**
HASANOĞLU A., OKUR İ.
in: TEMEL PEDİATRİ, HASANOĞLU ENVER, DÜŞÜNSEL RUHAN, BİDECİ AYSUN, BODUROĞLU KORAY, Editor, GÜNEŞ
TIP KİTABEVLERİ, Ankara, pp.30-44, 2020
- IX. **Pediyatrik Hastalıklara Özel Beslenme ve Diyet Yönetimi**
Okur İ. (Editor)
Akademisyen Kitabevi, Ankara, 2019
- X. **Dikkat Eksikliği Ve Hiperaktivite Bozukluğunda Nörometabolik Değerlendirme**
OKUR İ.
in: Dikkat Eksikliği Ve Hiperaktivite Bozukluğu, Şebnem Soysal, Editor, Nobel, pp.483-488, 2019
- XI. **Normal Çocuklukta Beslenme ve Beslenme Bozuklukları**
TÜMER L., İNCİ A., OKUR İ., Kasapkara Ç. S. , OLGAC M. A. B.
in: Lange - Current Tanı ve Tedavi Pediatri, Prof.Dr. Enver Hasanoğlu Prof.Dr. Aysun Bideci Prof.Dr. Elif N. Özmert
Prof.Dr. Sevcan A. BAKKALOĞLU EZGÜ, Editor, ema tıp kitapevi, pp.281-308, 2018
- XII. **Peroksizomal Bozukluklar**
OKUR İ.
in: Yurdakök Pediatri, Murat Yurdakök, Editor, Güneş Tıp Kitapevleri, Ankara, pp.1813-1820, 2017

Refereed Congress / Symposium Publications in Proceedings

- I. **Triamterene-induced suppression of R227X premature termination codon in Fabry disease**
Dundar H., Udgu B., BİBEROĞLU G., İnci A., Ezgu F., IŞIK GÖNÜL İ., OKUR İ., TÜMER L.
16th Annual Research Meeting of the WORLDSymposium(TM), Florida, United States Of America, 10 - 14 February
2020, vol.129
- II. **Screening of twelve lysosomal storage diseases with LC-MS/MS in Gazi university hospital in Turkey:
The first results of validation**
BİBEROĞLU G., İNCİ A., DERİN B., OKUR İ., EZGÜ F. S. , TÜMER L.
SSIEM, 3 - 06 September 2019
- III. **Beneficial Effects of Modified Atkins Diet in Glycogen Storage Disease Type IIIa**
OLGAC M. A. B. , İNCİ A., OKUR İ., Kasapkara Ç. S. , BİBEROĞLU G., OĞUZ A. D. , Aktaş E., EZGÜ F. S. , TÜMER L.
SSIEM 2019, 3-6th September, 2019, Rotterdam-The Netherlands, 3 - 06 September 2019
- IV. **Next generation DNA sequencing as an initial diagnostic method for congenital defects of
glycosylation**
EZGÜ F. S. , İNCİ A., Çiftçi B., TÜMER L., OKUR İ., Topçu B., Hasanoğlu A.
SSIEM 2019, 3-6th September, 2019, Rotterdam-The Netherlands, 3 - 06 September 2019
- V. **Diyet tedavisine cevap veren HMG-CoA liyaz enzim eksikliği olan iki olgu**
KOÇ N., KUYUCU A., GÜNDÜZ M., OKUR İ., ÖZAYDIN E.
Hacettepe Beslenme ve Diyetetik Günleri IV. Mezuniyet Sonrası Eğitim Kursu, Ankara, Turkey, 27 - 29 June 2019
- VI. **Cornelia de Lange Syndrome and Glycogen Storage Disease Together in a Patient**
KILIÇ A., EMECAN ŞANLI M., ÖZSAYDI E., İNCİ A., OKUR İ., TÜMER L., EZGÜ F. S.
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BİBEROĞLU G., İNCİ A., DERİN B., OKUR İ., EZGÜ F. S. , TÜMER L.
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- VIII. **Could Targeted Next Generation Sequencing Be A First Line Diagnostic Method for Lysosomal Storage Disease?**
CENGİZ F. B. , İNCİ A., BİBEROĞLU G., Çiftçi B., Topçu B., Tokgöz D., Yazar Ö. F. , Gökmenoğlu H., Raj Y., OKUR İ., et al.
International Inborn Errors of Metabolism and Nutrition Congress, 10 - 14 April 2019
- IX. **Familial Hyperphosphatemic Tumoral Calcinosis in an Unusual Site**
Emecan Şanlı M., Özsaydı E., kılıç m., İNCİ A., OKUR İ., EZGÜ F. S. , TÜMER L.
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- X. **Novel Mutation in Two Siblings with Normouricemic Lesch Nyhan Syndrome**
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- XI. **A Very Rare Disease: Hyperornithinemia-Hyperammonemia-Homocitrullinuria (Hhh) Syndrome**
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- XII. **Could Targeted Next Generation Sequencing Be A First Line Diagnostic Method for Lysosomal storage Diseases**
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- XIII. **Growth Hormone Treatment: Reverses Catabolic Process in Inborn Errors of Metabolism**
İNCİ A., OKUR İ., AKKUZU E., DÖĞER E., BİBEROĞLU G., KALKAN G., TÜMER L., EZGÜ F. S.
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- XIV. **Hyperinsulinemic Hypoglycemia: Think of GLUD1 Gene Mutation Leading To Hyperinsulinism/Hyperammonemia (HI/HA) Syndrome**
Emecan Şanlı M., kılıç m., Özsaydı E., İNCİ A., OKUR İ., TÜMER L., EZGÜ F. S.
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- XV. **Novel Mutation in FBP1 Gene Presenting with Recurrent Episodes of Vomiting in A Child**
Emecan Şanlı M., kılıç m., Özsaydı E., İNCİ A., OKUR İ., EZGÜ F. S. , TÜMER L.
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- XVI. **ICV-administered tralostinidase alfa (BMN 250 NAGLU-IGF2) is well-tolerated and reduces heparan sulfate accumulation in the CNS of subjects with Sanfilippo syndrome type B (MPS IIIB)**
Cleary M., Muschol N., Luz Couce M., Harmatz P., Lee J., Lin S., OKUR İ., Ezgu F., Peters H., Villarreal M. S. , et al.
15th Annual Research Meeting of the WORLDSymposium(TM), Florida, United States Of America, 4 - 07 February 2019, vol.126
- XVII. **Respiratory system involvement of 41 Mucopolysaccharidosis patients with the evaluation of KL-6, SPA and SPD levels**
İNCİ A., OKUR İ., Yılmaz Demirtaş C., BİBEROĞLU G., ASLAN A. T. , EZGÜ F. S. , TÜMER L.
15 th MEMG, Beirut, 29 November - 02 December 2018
- XVIII. **UNIQUE CLINICAL AND MOLECULAR FINDINGS IN LARGE COHORT OF PATIENTS WITH GAUCHER DISEASE FROM TURKEY**
Akay Tayfun G., OKUR İ., BİBEROĞLU G., TÜMER L., İNCİ A., Küçükcongür A., Hasanoğlu A., EZGÜ F. S.

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