

Prof. LEYLA TÜMER

Personal Information

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Education Information

Post Doctorate of Medicine, Gazi University, Sağlık Bilimleri Enstitüsü, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı, Turkey
1997 - Continues

Expertise In Medicine, Gazi University, Tıp Fakültesi, Turkey 1987 - 1993

Under Graduate, Ankara University, Tıp Fakültesi, Tıp Pr., Turkey 1981 - 1987

Research Areas

Health Sciences

Academic Titles / Tasks

Professor, Gazi University, Tıp Fakültesi, 2015 - Continues

Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2006 - Continues

Professor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 1998 - Continues

Advising Theses

TÜMER L., Gazi Üniversitesi çocuk beslenme ve metabolizma bölümünde takipli glikojen depo ve fenilketonüri hastalarının malnutrisyon durumunun değerlendirilmesi, Expertise In Medicine, C.MULUK(Student), 2020

TÜMER L., Effect of proton pump inhibitors on vitamin b12, iron, calcium and magnesium absorption, Expertise In Medicine, S.KANMAZ(Student), 2014

TÜMER L., Use of continuous glucose monitoring evaluating the accuracy and the effect on metabolic parameters in patients with gsd i followed at GaziUuniversity hospital pediatric metabolic unit, Expertise In Medicine, Ç.SEHER(Student), 2012

TÜMER L., İlkokul çağındaki çocuklarda A vitamini ve çinko düzeyinin belirlenmesi ve etki eden faktörlerin değerlendirilmesi, Expertise In Medicine, D.VURALLI(Student), 2010

TÜMER L., 1- 16 YAS ARASI ÇOCUKLARDA D VİTAMİNİ DÜZEYİ ve BUNA ETKİ EDEN FAKTÖRLERİN BELİRLENMESİ, A.ÖDEN(Student), 2010

TÜMER L., İLKOKUL ÇAĞINDAKİ ÇOCUKLARDA A VİTAMİNİ VE ÇİNKO DÜZEYİNİN BELİRLENMESİ VE ETKİ EDEN FAKTÖRLERİN DEĞERLENDİRİLMESİ, D.VURALLI(Student), 2010

TÜMER L., 0-16 yaş arası çocuklarda D vitamini yetersizliği ve buna etki eden faktörlerin belirlenmesi, Expertise In Medicine, A.ÖDEN(Student), 2009

TÜMER L., Glikojen depo tip 1A ve Glikojen depo tip 1B hastalarında sık gözlenen Glukoz-6- fosfataz ve Glukoz-6- fosfat taşıyıcı gen mutasyonlarının mikroelektronik array teknolojisi ile araştırılması, Post Doctorate of Medicine, F.TUBA(Student), 2009

TÜMER L., Okul çağı çocuklarında hiperlipidemi taraması, Expertise In Medicine, O.DERİNÖZ(Student), 2005

TÜMER L., Çocukluk çağı obezitesinde tümör nekrozis faktör-alfa-, plazminojen aktivatör inhibitör-1 ve adiponektin düzeyleri, Expertise In Medicine, F.ÖZBAY(Student), 2003

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

- I. **A CASE OF GLYCOGEN STORAGE DISEASE TYPE 1a MIMICKING FAMILIAL CHYLOMICRONEMIA SYNDROME**
Olgac A., OKUR İ., BİBEROĞLU G., EZGÜ F. S. , TÜMER L.
BALKAN JOURNAL OF MEDICAL GENETICS, vol.24, no.1, pp.103-105, 2021 (Journal Indexed in SCI)
- II. **Congenital defects of glycosylation: Novel presentations with mainly neurological involvement and variable dysmorphic features**
İNCİ A., Cengiz B., BİBEROĞLU G., OKUR İ., ARHAN E., ÖNER A. Y. , KASAPKARA Ç. S. , Kucukcongar A., TÜMER L., Ezgu F.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, 2021 (Journal Indexed in SCI)
- III. **The first case with FBXL4 mutation successfully treated with a parenteral ketogenic diet for lactic acidosis**
İNCİ A., Aktas E., Cengiz Ergin F. B. , OKUR İ., BİBEROĞLU G., EZGÜ F. S. , TÜMER L.
JOURNAL OF PARENTERAL AND ENTERAL NUTRITION, 2021 (Journal Indexed in SCI)
- IV. **Ultra-Rare Disorder in a Young Girl with Lipodystrophy: Analbuminemia**
İNCİ A., Arslan B., OKUR İ., BİBEROĞLU G., ŞANLI M. E. , ÖZSAYDI AKTAŞOĞLU E., KILIÇ A., TÜMER L., EZGÜ F. S.
INDIAN JOURNAL OF PEDIATRICS, 2021 (Journal Indexed in SCI)
- V. **Autism: Screening of inborn errors of metabolism and unexpected results**
İNCİ A., ÖZASLAN A., OKUR İ., BİBEROĞLU G., GÜNEY E., EZGÜ F. S. , TÜMER L., İŞERİ E.
AUTISM RESEARCH, 2021 (Journal Indexed in SCI)
- VI. **The chemical chaperone 4-phenylbutyrate enhances alpha-galactosidase activity subsequent to stop-codon read-through therapy with triamterene in Fabry R227X fibroblasts**
Dündar H., Biberoglu G., İnci A., Işık Gönül İ., Okur İ., Tümer L., Ezgü F. S.
MOLECULAR GENETICS AND METABOLISM, vol.132, no.2, 2021 (Journal Indexed in SCI)
- VII. **Familial hyperphosphatemic tumoral calcinosis in an unusual and usual sites and dramatic improvement with the treatment of acetazolamide, sevelamer and topical sodium thiosulfate**
ŞANLI M. E. , KILIÇ A., ÖZSAYDI AKTAŞOĞLU E., İNCİ A., OKUR İ., Ezgu F., TÜMER L.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.34, no.6, pp.813-816, 2021 (Journal Indexed in SCI)
- VIII. **Two patients from Turkey with a novel variant in the GM2A gene and review of the literature**
İNCİ A., ERGİN F. B. , BİBEROĞLU G., OKUR İ., EZGÜ F. S. , TÜMER L.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.34, no.6, pp.805-812, 2021 (Journal Indexed in SCI)
- IX. **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, no.4, pp.233-241, 2020 (Journal Indexed in SCI)
- X. **A new NBIA patient from Turkey with homozygous C19ORF12 mutation**
Kasapkara C. S. , TÜMER L., Gregory A., Ezgu F., İNCİ A., Derinkuyu B. E. , Fox R., Rogers C., Hayflick S.
ACTA NEUROLOGICA BELGICA, vol.119, no.4, pp.623-625, 2019 (Journal Indexed in SCI)
- XI. **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)
- XII. **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, no.5, pp.355-359, 2018 (Journal Indexed in SCI)
- XIII. **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, no.3, pp.243-245, 2018 (Journal Indexed in SCI)
- XIV. **A Myopathy, Lactic Acidosis, Sideroblastic Anemia (MLASA) Case Due to a Novel PUS1 Mutation**
Kasapkara C. S., TÜMER L., Zanetti N., Ezgu F., Lamantea E., Zeviani M.
TURKISH JOURNAL OF HEMATOLOGY, vol.34, no.4, pp.376-377, 2017 (Journal Indexed in SCI)
- XV. **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, no.3, pp.281-284, 2016 (Journal Indexed in SCI)
- XVI. **Secondary Hemophagocytosis in Propionic Acidemia**
Kasapkara C. S., Kangin M., Ozmen B. O., Ozbek M. N., Demir R., Karatas M., TÜMER L., EZGÜ F. S., Hasanoglu A.
IRANIAN JOURNAL OF PEDIATRICS, vol.25, no.3, 2015 (Journal Indexed in SCI)
- XVII. **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, no.3, pp.400-405, 2015 (Journal Indexed in SCI)
- XVIII. **Diagnostic Dilemma: Osteopetrosis with superimposed rickets causing Neonatal Hypocalcemia**
Olgac A., TÜMER L., Boyunaga O., Kizilkaya M., Hasanoglu A.
JOURNAL OF TROPICAL PEDIATRICS, vol.61, no.2, pp.146-150, 2015 (Journal Indexed in SCI)
- XIX. **BCS1L gene mutation causing GRACILE syndrome: case report**
Kasapkara C. S., TÜMER L., EZGÜ F. S., Kucukcongar A., Hasanoglu A.
RENAL FAILURE, vol.36, no.6, pp.953-954, 2014 (Journal Indexed in SCI)
- XX. **Vitamin A status and factors associated in healthy school-age children**
VURALLI KARAOĞLAN D., TÜMER L., Hasanoglu A., BİBEROĞLU G., PAŞAOĞLU H.
CLINICAL NUTRITION, vol.33, no.3, pp.509-512, 2014 (Journal Indexed in SCI)
- XXI. **Expanding the Mutational Spectrum of CRLF1 in Crisponi/CISS1 Syndrome**
Piras R., Chiappe F., La Torraca I., Buers I., Usala G., Angius A., Akin M. A., Basel-Vanagaite L., Benedicenti F., Chiodin E., et al.
HUMAN MUTATION, vol.35, no.4, pp.424-433, 2014 (Journal Indexed in SCI)
- XXII. **Home sleep study characteristics in patients with mucopolysaccharidosis**
Kasapkara C. S., TÜMER L., ASLAN A. T., Hasanoglu A., EZGÜ F. S., Kucukcongar A., Tunca Z., KÖKTÜRK O.
SLEEP AND BREATHING, vol.18, no.1, pp.143-149, 2014 (Journal Indexed in SCI)
- XXIII. **Diagnosis of glycine encephalopathy in a pediatric patient by detection of a GLDC mutation during initial next generation DNA sequencing**
Ezgu F., Ciftci B., Topcu B., Adiyaman G., Gokmenoglu H., Kucukcongar A., Kasapkara C., Biberoglu G., TÜMER L., Hasanoglu A.
METABOLIC BRAIN DISEASE, vol.29, no.1, pp.211-213, 2014 (Journal Indexed in SCI)
- XXIV. **DGUOK-Related Mitochondrial DNA Depletion Syndrome in a Child With an Early Diagnosis of Glycogen Storage Disease**
Kasapkara C. S., TÜMER L., Kuecukcongar A., Hasanoglu A., Seneca S., De Meirleir L.
JOURNAL OF PEDIATRIC GASTROENTEROLOGY AND NUTRITION, vol.57, no.5, 2013 (Journal Indexed in SCI)
- XXV. **Sleep study characteristics in patients with mucopolysaccharidosis**
ASLAN A. T., Kasapkara C., TÜMER L., Hasanoglu A., Ezgu F.
EUROPEAN RESPIRATORY JOURNAL, vol.42, 2013 (Journal Indexed in SCI)
- XXVI. **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, no.4, pp.428-433, 2013 (Journal Indexed in SCI)
- XXVII. **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)
- XXVIII. **Apheresis-inducible cytokine pattern change in children with homozygous familial hypercholesterolemia**
Kucukcongar A., Yenicesu I., TÜMER L., Kasapkara C. S. , EZGÜ F. S. , Pasaoglu O., Demirtas C., ÇELİK B., Dilsiz G., Hasanoglu A.
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- XXIX. **Rhabdomyolysis and acute kidney injury in two children: Questions**
Fidan K., Kandur Y., TÜMER L., Hasanoglu A., Soylemezoglu O.
PEDIATRIC NEPHROLOGY, vol.28, no.6, pp.899-902, 2013 (Journal Indexed in SCI)
- XXX. **Asymmetric dimethylarginine (ADMA) and L-arginine levels in children with glycogen storage disease type I**
Kasapkara C. S. , TÜMER L., BİBEROĞLU G., Kasapkara A., Hasanoglu A.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.26, pp.427-431, 2013 (Journal Indexed in SCI)
- XXXI. **AN EXTREMELY RARE CASE: OSTEOSCLEROTIC METAPHYSEAL DYSPLASIA**
Kasapkara C. S. , Kucukcongar A., BOYUNAGA Ö. L. , Bedir T., Oncu F., Hasanoglu A., TÜMER L.
GENETIC COUNSELING, vol.24, no.1, pp.69-74, 2013 (Journal Indexed in SCI)
- XXXII. **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, no.1, pp.35-37, 2012 (Journal Indexed in SCI)
- XXXIII. **Frequency of vitamin D insufficiency in healthy children between 1 and 16 years of age in Turkey**
Akman A. O. , TÜMER L., Hasanoglu A., Ilhan M., Cayci B.
PEDIATRICS INTERNATIONAL, vol.53, no.6, pp.968-973, 2011 (Journal Indexed in SCI)
- XXXIV. **Analysis of acylcarnitine levels by tandem mass spectrometry in epileptic children receiving valproate and oxcarbazepine**
CANSU A., SERDAROĞLU A., BİBEROĞLU G., TÜMER L., HIRFANOĞLU T., EZGÜ F. S. , Hasanoglu A.
EPILEPTIC DISORDERS, vol.13, no.4, pp.394-400, 2011 (Journal Indexed in SCI)
- XXXV. **A rare case of severe lactic acidosis in a preterm infant: lack of thiamine during total parenteral nutrition**
Oguz S. S. , Ergenekon E., TÜMER L., KOÇ E., Turan O., Onal E., TÜRKYILMAZ C., Atalay Y.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.24, pp.843-845, 2011 (Journal Indexed in SCI)
- XXXVI. **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, no.5, pp.522-527, 2011 (Journal Indexed in SCI)
- XXXVII. **Very long-chain acyl CoA dehydrogenase deficiency which was accepted as infanticide**
Eminoglu T. F. , TÜMER L., OKUR İ., EZGÜ F. S. , BİBEROĞLU G., Hasanoglu A.
FORENSIC SCIENCE INTERNATIONAL, vol.210, 2011 (Journal Indexed in SCI)
- XXXVIII. **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, no.6, pp.799-801, 2011 (Journal Indexed in SCI)
- XXXIX. **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, no.1, pp.225-231, 2011 (Journal Indexed in SCI)
- XL. **Mitochondrial neurogastrointestinal encephalomyopathy (MNGIE): case report with a new mutation**
Baris Z., Eminoglu T., DALGIÇ B., TÜMER L., Hasanoglu A.
EUROPEAN JOURNAL OF PEDIATRICS, vol.169, no.11, pp.1375-1378, 2010 (Journal Indexed in SCI)
- XLI. **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, no.4, pp.478-481, 2009 (Journal Indexed in SCI)

- XLII. 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, no.3, pp.235-240, 2008 (Journal Indexed in SCI)
- XLIII. Crisponi Syndrome: A New Case With Additional Features and New Mutation in CRLF1**
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AMERICAN JOURNAL OF MEDICAL GENETICS PART A, no.24, pp.3237-3239, 2008 (Journal Indexed in SCI)

Articles Published in Other Journals

- I. The Evaluation of Skeletal Manifestations in Patients with Gaucher Disease**
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JOURNAL OF PEDIATRIC RESEARCH, vol.8, no.3, pp.257-261, 2021 (Journal Indexed in ESCI)
- II. Citrullinemia with an Atypical Presentation: Paroxysmal Hypoventilation Attacks**
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JOURNAL OF PEDIATRIC NEUROSCIENCES, vol.13, no.2, pp.276-278, 2018 (Journal Indexed in ESCI)
- III. "Double Hit" Homozygous Mutations for Two Different Rare Inborn Errors of Metabolism: A Burden for Countries with High Prevalences of Consanguineous Marriages**
Olgac A., TÜMER L., Ceylaner S., BİBEROĞLU G., Hasanoglu A.
JOURNAL OF PEDIATRIC RESEARCH, vol.5, no.1, pp.47-50, 2018 (Journal Indexed in ESCI)
- IV. 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, no.1, pp.1-5, 2017 (Journal Indexed in ESCI)
- V. L carnitine L propionyl carnitine and malondialdehyde levels of pediatric patients with solid tumor**
OKUR A., HASANOĞLU A., OĞUZ A., BİBEROĞLU G., ERTEM U., TÜMER L.
JOURNAL OF PEDIATRIC SCIENCES, vol.4, no.3, 2012 (Other Refereed National Journals)

Books & Book Chapters

- I. Normal Çocuklukta Beslenme ve Beslenme Bozuklukları**
TÜMER L., İNCİ A., OKUR İ., Kasapkar Ç. S. , OLGAÇ M. A. B.
in: Lange - Current Tanı ve Tedavi Pediatri, Prof.Dr. Enver Hasanoglu Prof.Dr. Aysun Bideci Prof.Dr. Elif N. Özmert Prof.Dr. Sevcin A. BAKKALOĞLU EZGÜ, Editor, ema tıp kitapevi, pp.281-308, 2018
- II. Mitokondriyal Hastalıklar**
TÜMER L., İNCİ A.
in: Yurdakök Pediatri, Murat Yurdakök, Editor, Güneş Tıp Kitapevi, Ankara, pp.1779-1790, 2017
- III. Yoğurt ve Laktoz İntoleransı**
TÜMER L., OLGAÇ M. A. B.
in: YoğurtLezzetin ve Sağlığın Öyküsü, Sevinç Yücecan, Editor, Matsis Matbaa, İstanbul, pp.107-1114, 2015

Refereed Congress / Symposium Publications in Proceedings

- I. Triamterene-induced suppression of R227X premature termination codon in Fabry disease**
Dündar H., Udgu B., Biberoglu G., Inci A., Ezgu F., Işık 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. **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.
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- IV. **Beneficial Effects of Modified Atkins Diet in Glycogen Storage Disease Type IIIa**
OLGAÇ M. A. B. , İNCİ A., OKUR İ., Kasapkara Ç. S. , BİBEROĞLU G., OĞUZ A. D. , Aktaş E., EZGÜ F. S. , TÜMER L.
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- V. **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.
International Inborn Errors Of Metabolism And Nutrition Congress, İstanbul, Turkey, 10 - 14 April 2019
- VI. **Screening of Twelve Lysosomal Storage Diseases with LC-MS/MS in Gazi University Hospital: The First Results of Validation.**
BİBEROĞLU G., İNCİ A., DERİN B., OKUR İ., EZGÜ F. S. , TÜMER L.
INTERNATIONAL INBORN ERRORS OF METABOLISM AND NUTRITION CONGRESS10 - 14 April 2019 Istanbul-Turkey, 10 - 14 April 2019
- VII. **A Novel Rars2 Mutation in Two Siblings with Microcephaly, Seizures and Liver Involvement**
EMİNOĞLU F. T. , Sevinç S., Karaköse Gök T., EZGÜ F. S. , İNCİ A., TÜMER L.
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- VIII. **Familial Hyperphosphatemic Tumoral Calcinosis in an Unusual Site**
Emecan Şanlı M., Özsaydı E., kılıç m., İNCİ A., OKUR İ., EZGÜ F. S. , TÜMER L.
International Inborn Errors Of Metabolism And Nutrition Congress 10 - 14 April 2019, Istanbul-Turkey, 10 - 14 April 2019
- IX. **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.
International Inborn Errors Of Metabolism And Nutrition Congress 10 - 14 April 2019, Istanbul-Turkey, 10 - 14 April 2019
- X. **A Very Rare Disease: Hyperornithinemia-Hyperammonemia-Homocitrullinuria (Hhh) Syndrome**
Özsaydı E., Emecan Şanlı M., kılıç m., İNCİ A., OKUR İ., TÜMER L., EZGÜ F. S.
International Inborn Errors Of Metabolism And Nutrition Congress 10 - 14 April 2019, Istanbul-Turkey, 10 - 14 April 2019
- XI. **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.
International Inborn Errors Of Metabolism And Nutrition Congress 10 - 14 April 2019, Istanbul-Turkey, 10 - 14 April 2019
- XII. **Could Targeted Next Generation Sequencing Be A First Line Diagnostic Method for Lysosomal storage Diseases**
İNCİ A., OKUR İ., AKKUZU E., DÖĞER E., BİBEROĞLU G., KALKAN G., TÜMER L., EZGÜ F. S.
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