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Education Information

Undergraduate Minor, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 2014 - 2017

Academic Titles / Tasks

Lecturer, Gazi University, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2018 - Continues

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

- I. **Congenital defects of glycosylation: Novel presentations with mainly neurological involvement and variable dysmorphic features**
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- II. **Ultra-Rare Disorder in a Young Girl with Lipodystrophy: Analbuminemia**
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- VIII. **Patient With Niemann-Pick Type C Presenting With a Jaw Mass Characterized With Lymph Node Involvement by Niemann-Pick Cells**
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Articles Published in Other Journals

- I. **Citrullinemia with an Atypical Presentation: Paroxysmal Hypoventilation Attacks**
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- I. **Yağ Asidi Oksidasyon Bozuklukları**
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- II. **Normal Çocuklukta Beslenme ve Beslenme Bozuklukları**
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- III. **Mitokondriyal Hastalıklar**
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in: Yurdakök Pediatri, Murat Yurdakök, Editor, Güneş Tıp Kitapevi, Ankara, pp.1552-1556, 2017

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- I. **Screening of twelve lysosomal storage diseases with LC-MS/MS in Gazi university hospital in Turkey: The first results of validation**
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- V. **Cornelia de Lange Syndrome and Glycogen Storage Disease Together in a Patient**
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- VI. **A Very Rare Disease: Hyperornithinemia-Hyperammonemia-Homocitrullinuria (Hhh) Syndrome**
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X. Novel Mutation in Two Siblings with Normouricemic Lesch Nyhan Syndrome

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XI. A Novel Rars2 Mutation in Two Siblings with Microcephaly, Seizures and Liver Involvement

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XV. 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.

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XVI. UNIQUE CLINICAL AND MOLECULAR FINDINGS IN LARGE COHORT OF PATIENTS WITH GAUCHER DISEASE FROM TURKEY

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- XXII. **An early diagnosis cerebretendinous xanthomatosis in a patient at the age of 15 years**
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- XXVIII. **Screening ALPL Gene Differences byNext Generation Sequence Techonology inPatients Having Low ALP Levels**
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- XXIX. **DiagnosticCapability ofNextGenerationDNA Sequencing With A 450 Gene Panel forInborn Errors of Metabolism**
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- XXXIII. **Evaluation of chitotriosidase and high sensitive c reactive protein levels in mucopolysaccharidosis**
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- XXXVIII. **Early initiation of investigational enzyme replacement therapy in a nine month old infant with mucopolysaccharidosis type VII**
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- XLIV. **Type 1 hypersensitivity reaction and desensitization with Elosulphase alpha**
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- XLVII. **Plasma acylcarnitine levels Are there New İnflammatory markers in lysosomal storage disease**
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- XLVIII. Is there any effect of acylcarnitines on proinflammatory process in obese children**
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- XLIX. Patient with Niemann Pick type C presenting with lymphatic involvement with Niemann Pick cells in the left jaw**
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- L. Mucopolysaccharidosis Type VII at an Early Age A good candidate for investigational enzyme replacement therapy**
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- LIII. Sol çenede Lenfatik tutulum ile giden Niemann Pick tip C olgusu**
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Citations

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