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Education Information

Doctorate, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , Turkey 2016 - 2021

Foreign Languages

English, B1 Intermediate

Certificates, Courses and Trainings

Personal Evolution, Eğiticinin Eğitimi, İstanbul Üniversitesi, 2020

Dissertations

Doctorate, SANTRAL ERKEN PUBERTE OLGULARINDA YENİ NESİL DİZİLEME İLE MOLEKÜLER PATOLOJİNİN AYDINLATILMASI, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 2021
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Postgraduate, Miks Lenfosist Kültür Testinin Allojeneik Kemik İliği Kök Hücre Naklinde Hasta-Verici Parametreleriyle İlişkisi, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 2006

Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Medical Genetics

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II. **Clinical and Molecular Findings of Nine Cases with Tay- Sachs Disease From Turkiye**
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III. **CLINICAL AND MOLECULAR RESULTS OF SIX CASES WITH ROBERTS SYNDROME: REVIEW OF CASES FROM TURKIYE**
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I. **Çoklu Bağlanmaya Bağımlı Prob Amplifikasyonu (Multıplex Lıgatıondependent Probe Amplıfıcatıon – MLPA)**
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in: Pediatri Pratiğinde Genetik Testlerin Seçimi ve Yorumlanması, Prof. Dr. Beyhan Tüysüz, Editor, Nobel Yayınevi, Ankara, pp.102-111, 2023

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I. **Evaluation of metabolic syndrome risk using metabolic syndrome z-score in Bardet-Biedl Syndrome patients with various genotypes**
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62nd Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), Liverpool, England, 16 - 18 November 2024, pp.385

II. **Unraveling the complexity of Phelan-McDermid syndrome: A multifaceted exploration of clinical and molecular/cytogenetic findings in four cases**
Yücesoy M. A., Akbaş S., Memiş G., Konur E., Durmaz D., Karaman V., Şentürk L., Altunoğlu U., Şengenç E., Maraş Genç H., et al.
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III. **The complexity of Phelan-McDermid syndrome: A multifaceted exploration of clinical and molecular/cytogenetic findings in four cases.**
Yücesoy M. A., Akbaş S., Memiş G., Konur Akbaş E. N., Durmaz D., Karaman V., Şentürk L., Altunoğlu U., Şengenç E., Maraş Genç H., et al.
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IV. **Aarskog syndrome: Expanding the phenotypic and molecular spectrum through a new case series including two adult patients (POSTER ID EP13.093)**
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V. **Gross deletion in KIF11: A de novo occurrence**
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