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Published journal articles indexed by SCI, SSCI, and AHCI

- I. **First note on marine-like cementation of Late Holocene beachrock, Iznik Lake (Turkey)**
ERGİNAL A. E., Kiyak N. G., ÖZTÜRK M., Yiğitbaş E., Bozcu M., Avcıoğlu M., ÖZTÜRK B.
GEOCHRONOMETRIA, vol.39, no.1, pp.76-83, 2012 (SCI-Expanded)

Articles Published in Other Journals

- I. **Amyand's hernia detected incidentally in two patients**
Resorlu M., Aylanç N., KARATAĞ O., ÖZTÜRK M.
BMJ Case Reports, vol.2017, 2017 (Scopus)

Books & Book Chapters

- I. **OTOİNFLAMATUAR HASTALIKLAR VE GENETİK**
ÖZTÜRK M., ALBUZ B., SILAN F.
in: Türkiye Klinikleri Tıbbi Genetik - Özel Konular Genetik ve Multidisipliner Yaklaşımlar, Prof Dr C. Nur SEMERCİ
GÜNDÜZ, Editor, <http://www.turkiyeklinikleri.com>, Ankara, pp.49-61, 2019

Refereed Congress / Symposium Publications in Proceedings

- I. **BCL11B gene may be a candidate gene for mastocytosis in a patient with partial trisomy of distal 14q**
ALBUZ B., SILAN F., ÖZTÜRK M., YILDIZ O., ÖZDEMİR Ö.
13th Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.22, pp.156-1156
- II. **A rare heterochromatin polymorphism of chromosome 6 associated with recurrent miscarriage: A case report**
ALBUZ B., SILAN F., ÖZTÜRK M., KARAKAYA T., ÖZDEMİR Ö.
13.Balkan Congress of Human Genetics, Edirne, Turkey, 17 - 20 April 2019, vol.0
- III. **Fetus with partial trisomy 4 and t(216) due to maternal complex rearrangement involving three chromosomes: a case report**
ALBUZ B., ÇETİN G. O., ÖZTÜRK M., CANER V., SEMERCİ GÜNDÜZ C. N., BAĞCI G.
European Human Genetics Conference 2018, 16 - 19 June 2018
- IV. **A novel nonsense mutation in the EYA1 gene found in a patient with BOR syndrome**
ÖZTÜRK M., ZEYBEK S., DURSUN B., ALBUZ B., BOZ Ö., GÜRKAN H., ÇETİN G. O.
European Human Genetics Conference 2018, 16 - 19 June 2018

- V. **Further delineation of ACPHD syndrome and a novel mutation in DNAJC3**
ZEYBEK S., FARRE G., ÖZTÜRK M., ALBUZ B., BELTRAN S., BOWCOCK A., ÇETİN G. O.
European Human Genetics Conference 2018, 16 - 19 June 2018
- VI. **The co-existence of Nablus Mask-Like Syndrome and Klinefelter Syndrome**
BOZ Ö., SARIKEPE B., ZEYBEK S., ÖZTÜRK M., BAĞCI G., ÇETİN G. O.
European Human Genetics Conference 2018, 16 - 19 June 2018
- VII. **The co-existence of Nablus mask-like facial syndrome and klinefelter syndrome**
BOZ Ö., SARIKEPE B., ZEYBEK S., ÖZTÜRK M., BAĞCI G., ÇETİN G. O.
Erciyes Tıp Genetik Günleri 2018, Kayseri, Turkey, 7 - 10 March 2018, vol.40, pp.71
- VIII. **fetus with partial trisomy 4 and t (216) due to maternal complex rearrangement involving three chromosome: a case report**
ALBUZ B., ÇETİN G. O., ÖZTÜRK M., CANER V., SEMERCİ GÜNDÜZ C. N., BAĞCI G.
ERCİYES MEDICAL GENETICS DAYS 1017, Kayseri, Turkey, 11 - 13 May 2017
- IX. **unbalanced chromosomal rearrangement in fetus with congenital anomaly:a case report**
ÖZTÜRK M., ÇETİN G. O., ALBUZ B., CANER V., SEMERCİ GÜNDÜZ C. N., BAĞCI G.
ERCİYES MEDICAL GENETICS DAYS 2017, Kayseri, Turkey, 11 - 13 May 2017
- X. **A case with vascular anomalies: Differential diagnosis and management**
ALBUZ B., KOCAMAZ H., ÖZTÜRK M., SARIKEPE B., TEPELİ E., BAĞCI G., SEMERCİ GÜNDÜZ C. N.
ERCİYES MEDICAL GENETICS DAYS 2017, Kayseri, Turkey, 11 - 13 May 2017

Metrics

Publication: 14

Citation (WoS): 4

Citation (Scopus): 4

H-Index (WoS): 1

H-Index (Scopus): 1