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Research Areas

Medical Biology, Medical Genetics, Molecular Biology and Genetics, Genomics, Health Sciences

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Biallelic loss-of-function variants in EXOC6B are associated with impaired primary ciliogenesis and cause spondylo-epi-metaphyseal dysplasia with joint laxity type 3**
ŞİMŞEK KİPER P. Ö., Jacob P., Upadhyai P., TAŞKIRAN Z. E., Guleria V. S., KARAOSMANOĞLU B., İMREN G., GÖÇMEN R., Bhavani G. S., Kausthubham N., et al.
HUMAN MUTATION, vol.43, no.12, pp.2116-2129, 2022 (SCI-Expanded)

II. **Expanding the phenotypic spectrum of TNFRSF11A-associated dysosteosclerosis: a case with intracranial extramedullary hematopoiesis**
Xue J., Simsek-Kiper P. O., ÜTİNE G. E., Yan L., Wang Z., Taskiran E. Z., KARAOSMANOĞLU B., İMREN G., GÖÇMEN R., Nishimura G., et al.
JOURNAL OF HUMAN GENETICS, vol.66, no.6, pp.607-611, 2021 (SCI-Expanded)

Refereed Congress / Symposium Publications in Proceedings

I. **Allel-Spesifik Anti-Sense Oligonükleotidler: Kalıtsal Retina Hastalıklarında ''Ince Ayar'' Tedavi Mümkün Olabilir Mi?**
Karaosmanoğlu B., İmren G., Ütine G. E., Taylan-Şekeroğlu H., Taşkıran Z. E.
15. Ulusal Tıbbi Genetik Kongresi, Muğla, Turkey, 9 - 13 November 2022

II. **Büyük veriden hücre karakterizasyonuna: Transkripsiyon faktörü kataloglarınınoluşturulması**
İMREN G., KARAOSMANOĞLU B., ÇELEBİ SALTIK B., ESENDAĞLI G., TAŞKIRAN Z. E.
Tıbbi Biyoloji ve Genetik Kongresi, Turkey, 27 - 30 October 2019

Metrics

Publication: 5
Citation (WoS): 5
Citation (Scopus): 5
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