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Doctorate, Hacettepe University, Sağlık Bilimleri Enstitüsü, Genetik, Turkey 1998 - 2002
Expertise In Medicine, Hacettepe University, Tıp Fakültesi, Pediatri, Turkey 1988 - 1993
Undergraduate, Hacettepe University, Tıp Fakültesi, Turkey 1982 - 1988

Foreign Languages

English, C1 Advanced

Dissertations

Doctorate, Nöral tüp defektlerinin etiyolojisinde 5, 10 metilentetrahidrofolat redüktaz gen polimorfizmlerinin rolü, Hacettepe Üniversitesi, Sağlık Bilimleri Fakültesi, 2002

Research Areas

Medicine, Internal Medicine Sciences, Child Health and Diseases, Medical Genetics, Health Sciences

Academic Titles / Tasks

Professor, Hacettepe University, Tıp Fakültesi (İngilizce), Dahili Tıp Bilimleri Bölümü, 1995 - Continues

Academic and Administrative Experience

Hacettepe Üniversitesi, Çocuk Sağlığı Enstitüsü, 2007 - Continues
Hacettepe Üniversitesi, Hacettepe Üniversitesi Hastaneleri Araştırma Ve Uygulama Merkezi, 2008 - 2010
Hacettepe Üniversitesi, Hacettepe Üniversitesi Hastaneleri Araştırma Ve Uygulama Merkezi, 2000 - 2008

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Hyaline fibromatosis syndrome: a rare, yet recognizable syndrome**
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XXIV. **Natural history of TRPV4-Related disorders: From skeletal dysplasia to neuromuscular phenotype**
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EUROPEAN JOURNAL OF PAEDIATRIC NEUROLOGY, pp.46-55, 2021 (SCI-Expanded)

XXV. **Genetic disorders with symptoms mimicking rheumatologic diseases: A single-center retrospective study**
KAYA AKCA Ü., ŞİMŞEK KİPER P. Ö., ÜREL DEMİR G., SAĞ E., ATALAY E., ÜTİNE G. E., ALİKAŞİFOĞLU M., BODUROĞLU O. K., BİLGİNER Y., ÖZEN S.
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XXVI. **Two Siblings with Kaufman Oculocerebrofacial Syndrome Resembling Oculoauriculovertebral Spectrum**
ÜREL DEMİR G., Aydin B., KARAOSMANOĞLU B., AKGÜN DOĞAN Ö., Taskiran E. Z., ŞİMŞEK KİPER P. Ö., ÜTİNE G. E.,

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XXIX. **Further Phenotypic Delineation of Partial Trisomy 17q and Partial Monosomy 20q due to Rare t(17;20)**
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XXXIV. **Hyperphosphatasia with mental retardation syndrome type 4 In two siblings-expanding the phenotypic and mutational spectrum**
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EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.62, no.6, 2019 (SCI-Expanded)

XXXV. **An eight-case 1q21 region series: novel aberrations and clinical variability with new features**
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XXXVII. **Effects of Vitamin D and estrogen receptor polymorphisms on bone mineral density in adolescents with anorexia nervosa**
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Simsek-Kiper P. O., KOŞUKCU C., Akgun-Dogan O., GÖÇMEN R., ÜTİNE G. E., SOYER T., Korkmaz-Toygar A., Nishimura G., ALİKAŞİFOĞLU M., Boduroglu K.

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Simsek-Kiper P., Taskiran E., Kosukcu C., Urel-Demir G., Akgun-Dogan O., Yilmaz G., Utine G., Nishimura G., Boduroglu K., Alikasifoglu M.
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Metrics

Publication: 209
Citation (WoS): 1163
Citation (Scopus): 1299
H-Index (WoS): 19
H-Index (Scopus): 20