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Education Information

Doctorate, Hacettepe University, Çocuk Sağlığı Enstitüsü, Moleküler Metabolizma, Turkey 2016 - 2016
Postgraduate, Hacettepe University, Sağlık Bilimleri Enstitüsü, Biyoinformatik, Turkey 2013 - 2016
Undergraduate, Hacettepe University, Fen Fakültesi, Biyoloji, Turkey 2009 - 2013

Foreign Languages

English, C1 Advanced

Dissertations

Postgraduate, Otozomal dominant peters anomalisi tanısı alan bir ailede yeni gen araştırması, Hacettepe Üniversitesi, Sağlık Bilimleri Enstitüsü, Biyoinformatik (Yl) (Tezli), 2015

Research Areas

Medicine, Health Sciences, Fundamental Medical Sciences, Biostatistics and Medical Informatics, Medical Biology

Academic Titles / Tasks

Research Assistant, Hacettepe University, Sağlık Bilimleri Enstitüsü, Biyoinformatik A.B.D., 2016 - Continues

Published journal articles indexed by SCI, SSCI, and AHCI

I. **Implication of transcription factor FOXD2 dysfunction in syndromic congenital anomalies of the kidney and urinary tract (CAKUT)**
Riedhammer K. M., Nguyen T. T., KOŞUKCU C., Calzada-Wack J., Li Y., Assia Batzir N., SAYGILI S. K., Wimmers V., Kim

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V. **One Disease with two Faces: Semidominant Inheritance of a Novel HTRA1 Mutation in a Consanguineous Family**
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VIII. **Frequency of HLA Class I and Class II Alleles in Patients with CVID from Turkey**
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X. **Atypical Presentation of Sengers Syndrome: A Novel Mutation Revealed with Postmortem Genetic Testing.**
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Kilic M., Erguner B., KOŞUKCU C., ÖZGÜL R. K.
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XII. **Further expanding the mutational spectrum and investigation of genotype-phenotype correlation in 3M syndrome**
Simsek-Kiper P. O., Taskiran E., KOŞUKCU C., ARSLAN U. E., Cormier-Daire V., Gonc N., Ozon A., ALİKAŞİFOĞLU A., KANDEMİR N., ÜTİNE G. E., et al.
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XVII. **Further delineation of spondyloepimetaphyseal dysplasia Faden-Alkuraya type: A RSPRY1-associated spondylo-epi-metaphyseal dysplasia with cono-brachydactyly and craniosynostosis**
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XX. **Clinical, demographic and nosologic characterisation of the genetic disorders of the skeleton in Turkey: The Skeletal Dysplasia registry**
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XXIV. **The effects of larval diet restriction on developmental time, preadult survival, and wing length in Drosophila melanogaster**
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TURKISH JOURNAL OF ZOOLOGY, vol.39, no.3, pp.395-403, 2015 (SCI-Expanded)

XXV. **Mutations In Anks6 Cause A Nephronophthisis-like Phenotype With End Stage Renal Disease**
Taskiran E., Korkmaz E., Gucer S., KOŞUKCU C., Kaymaz F., Koyunlar C., Bryda E. C., Chaki M., Lu D. D., Vadnagara K.,

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Taskiran E. Z., Korkmaz E., Gucer S., KOŞUKCU C., Kaymaz F., Koyunlar C., Bryda E. C., Chaki M., Lu D., Vadnagara K., et al.
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I. **Current Diagnostic Methods in Primary Ciliary Dyskinesia: Hacettepe University Experience**
EMİRALİOĞLU N., TAŞKIRAN Z. E., KOŞUKCU C., BİLGİÇ E., ATİLLA P., KAYA Z. B., GÜNAYDIN R. Ö., yüzbaşoğlu a., Ademhan D., Eryılmaz Polat S., et al.
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Refereed Congress / Symposium Publications in Proceedings

I. **Sendromik olmayan zihinsel yetersizlikte tüm ekzom dizilemenin tanısal verimi**
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16. Tıbbi Biyoloji ve Genetik Kongresi, Turkey, 25 - 27 October 2019

II. **The Skeletal Dysplasia Registry: Hacettepe Experience**
ŞİMŞEK KİPER P. Ö., TAŞKIRAN Z. E., KOŞUKCU C., ARSLAN U. E., ÜTİNE G. E., ALANAY Y., ALİKAŞİFOĞLU M., BODUROĞLU O. K.
The 14th biannual International Skeletal Dysplasia Society Meeting, Oslo, Norway, 11 - 14 September 2019

III. **Severe Motor Mental Retardation with Microcepahlyand Hypomyelination due to PYCR2 Gene variant in aLarge Family**
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IV. **The Correlation Between Clinical Characteristics and Molecular Genetic Analysis Results of Primary Ciliary Dyskinesia Patients: Hacettepe University Experience**
EMİRALİOĞLU N., Taskiran E., KOŞUKCU C., BİLGİÇ E., ATİLLA P., Kaya B., Gunaydin O., YÜZBAŞIOĞLU A., Tugcu G. D., Ademhan D., et al.
International Congress of the European-Respiratory-Society (ERS), Madrid, Spain, 28 September - 02 October 2019, vol.54

V. **TIP III HEREDITER ANJIYOÖDEM ÖN TANILI HASTALARDA YENI NESIL DIZILEME ILE YENI GENLERIN ARAŞTIRILMASI: ÖN VERILER**
Koşukcu C., Uysal Soyer Ö., Karakaya G., Şahiner Ü. M., Damadoğlu E., Ocak M., Kaya S. B., Çakmak M., Kalyoncu A. F., Şekerel B. E., et al.
XXV. Ulusal Allerji ve Klinik İmmünoloji Kongresi, Antalya, Turkey, 17 - 21 November 2018

VI. **A Rare Cause of Hyperinsulinemic Hypoglycemia: Costello Syndrome**
VURALLI KARAOĞLAN D., KOŞUKCU C., TAŞKIRAN Z. E., ŞİMŞEK KİPER P. Ö., UTİNE G. E., BODUROĞLU O. K., ALİKAŞİFOĞLU A., ALİKAŞİFOĞLU M.
57th Annual ESPE European Society for Paediatric Endocrinology, Atina, Greece, 27 - 29 September 2018, vol.90, pp.351-352

VII. **Clinical, demographic and nosologic characterization of the genetic disorders of the skeleton in Turkey: The skeletal dysplasia registry.**
ŞİMŞEK KİPER P. Ö., ÜTİNE G. E., TAŞKIRAN Z. E., KOŞUKCU C., ARSLAN U. E., ALANAY Y., ALİKAŞİFOĞLU M.,

BODUROĞLU O. K.

European Human Genetics Conference, Milan, Italy, June 16-19, 2018., Milan, Italy, 16 - 19 June 2018

VIII. **PGAP3 geninde yeni tanımlanmış mutasyona bağlı hiperfosfatazya mental retardasyon sendromu**
AKGÜN DOĞAN Ö., ÜREL DEMİR G., KOŞUKCU C., TAŞKIRAN Z. E., ŞİMŞEK KİPER P. Ö., ÜTİNE G. E., ALİKAŞİFOĞLU M., BODUROĞLU O. K.
3. Ulusal Çocuk Genetik Sempozyumu, Antalya, Turkey, 11 - 13 October 2017

IX. **Further delineation of spondyloepimetaphyseal dysplasia Faden-Alkuraya type: a RSPRY1-associated spondylo-epi-metaphyseal dysplasia with cono-brachydactyly and craniosynostosis.**
ŞİMŞEK KİPER P. Ö., TAŞKIRAN Z. E., KOŞUKCU C., AKGÜN DOĞAN Ö., ÜTİNE G. E., BODUROĞLU O. K., ALİKAŞİFOĞLU M.
International Skeletal Dysplasia Society Meeting, 20 - 23 September 2017

X. **Identification of Molecular Pathology of Peters' Anomaly Segregating in a Large Autosomal Dominant Family**
KOŞUKCU C., ASLI K., ALANAY Y., KAVAK P., BERKER N., TAŞKIRAN Z. E., ALİKAŞİFOĞLU M., SEZERMAN O. U., AKARSU A. N.
10th International Symposium on Health Informatics and Bioinformatics (HIBIT 2017), KALKANLI, GUZELYURT, Cyprus (Kktc), 29 - 30 June 2017

XI. **RSPRY1 associated skeletal dysplasia: Spondylo-epi-metaphyseal dysplasia with conobrachydactyly and craniosynostosis**
ŞİMŞEK KİPER P. Ö., TAŞKIRAN Z. E., KOŞUKCU C., AKGÜN DOĞAN Ö., YILMAZ G., ÜTİNE G. E., Nishimura G., BODUROĞLU O. K., ALİKAŞİFOĞLU M.
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Supported Projects

BİRBEN E., UYSAL SOYER Ö., ŞEKEREL B. E., ŞAHİNER Ü. M., KOŞUKCU C., Project Supported by Higher Education Institutions, Herediter Anjioödem Tip III Tanılı Hastalarda Yeni Aday Genlerin Belirlenmesi, 2017 - 2022

ÜNAL D., ÜTİNE G. E., ÜNAL M. F., KOŞUKCU C., ŞİMŞEK KİPER P. Ö., ALİKAŞİFOĞLU M., Project Supported by Higher Education Institutions, İkiz Otistik Olgularda Genetik Farklılıklar, 2018 - 2021

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Metrics

Publication: 41
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