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Education Information

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

Foreign Languages

English, B2 Upper Intermediate

Dissertations

Doctorate, SANTRAL ERKEN PUBERTE OLGULARINDA YENİ NESİL DİZİLEME İLE MOLEKÜLER PATOLOJİNİN AYDINLATILMASI, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 2021
Postgraduate, SENDROMİK VE NON-SENDROMİK KRANİYOSİNOSTOZ OLGULARINDA FGFR1, FGFR2, FGFR3, TWIST1, MSX2, POR, FREM1 VE RAB23 GENLERİNDE MOLEKÜLER ANALİZLER, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 2015
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

Published journal articles indexed by SCI, SSCI, and AHCI

I. **A Rare Inherited Bone Marrow Failure Syndrome Disclosed by Reanalysis of the Exome Data of a Patient Evaluated for Cytopenia and Dysmorphic Features.**
Durmaz D., Aslanger A. D., Yavas Abali Z., Yilmaz Y., Karaman V., Yesil Sayin G., Toksoy G., Unuvar A., Uyguner Z. O.
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II. **Novel variants ensued genomic imprinting in familial central precocious puberty.**
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V. **Fibular Agenesis and Ball-Like Toes Mimicking Preaxial Polydactyly: Prenatal Presentation of du Pan Syndrome**
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VI. **Prenatal ultrasonographic features in Blomstrand osteochondrodysplasia: Antenatal case series confirmed by postmortem radiology and molecular diagnosis**
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Yılmaz Gulec E., Turgut G. T., Gezdırıcı A., Karaman V., Ozturk F. N., Avcı S., Kalaycı T., Senturk L., Ayaz A., Kayserılı H., et al.
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Ozturk A. P., Aslanger A., Ozturan E. K., Konur E. N., Gulec C., Karaman V., Yildiz M., Yesil G., Toksoy G., Poyrazoglu S., et al.
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XIII. **Mutations in AR or SRD5A2 Genes: Clinical Findings, Endocrine Pitfalls, and Genetic Features of Children with 46,XY DSD.**
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XV. **Sequence of MKRN3 and DLK1 genes in cases with familial central precocious puberty**
Karaman V., Karakilic-Ozturan E., Bas F., Poyrazoglu S., Basaran S., Darendeliler F., Uyguner Z. O.
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XVI. **Skeletal and molecular findings in 51 Cleidocranial dysplasia patients from Turkey**
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XVII. **Two cases with central precocious puberty caused by paternally inherited novel variants in DLK1gene**
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XVIII. **Prevalence, clinical characteristics and long-term outcomes of classical 11 β-hydroxylase deficiency (11BOHD) in Turkish population and novel mutations in CYP11B1 gene.**
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Annals of allergy, asthma & immunology : official publication of the American College of Allergy, Asthma, & Immunology, vol.114, no.2, pp.154-6, 2015 (SCI-Expanded)

XX. **NOVEL NLRP7 MUTATIONS IN FAMILIAL RECURRENT HYDATIDIFORM MOLES: ARE NLRP7 MUTATIONS AT THE SAME TIME A RISK FOR RECURRENT REPRODUCTIVE WASTAGE?**
Ulker V., Gurkan H., Tozkir H., Karaman V., Ozgur H., Numanoglu C., Gedikbasi A., Akbayir O., Uyguner Z. O.
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XXI. **Novel NLRP7 mutations in familial recurrent hydatidiform mole: are NLRP7 mutations a risk for recurrent reproductive wastage?**
Ülker V., Gurkan H., Tozkir H., Karaman V., Ozgur H., Numanoğlu C., Gedikbaşı A., Akbayır O., Uyguner Z. O.
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I. **JAG1 MUTATION SPECTRUM IN CASES WITH ALAGILLE SYNDROME FROM TURKIYE**
Aslanger A. D., Yildirim B. T., Kalayci T., Şentürk L., Avci Ş., Altunoğlu U., Güleç Ç., Karaman V., Doğan G., Önal Z., et al.
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II. **Clinical and Molecular Findings of Nine Cases with Tay- Sachs Disease From Turkiye**
ASLANGER A. D., GÜLEÇ Ç., KALAYCI T., Sengenc E., Avci S., Altunoglu U., KARAMAN V., TOKSOY G., KARACA M., Iscan A., et al.
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III. **CLINICAL AND MOLECULAR RESULTS OF SIX CASES WITH ROBERTS SYNDROME: REVIEW OF CASES FROM TURKIYE**
Aslanger A. D., Kalaycı T., Konur E. N., Güleç Ç., Avcı Ş., Altunoğlu U., Karaman V., Toksoy G., Karaman B., Başaran S., et al.
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IV. **GJB2-RELATED NON-SYNDROMIC HEARING LOSS VARIANTS' SPECTRUM AND THEIR FREQUENCY IN**

TURKISH POPULATION
Gulec C., Aslanger A. D., Karaman V., Wollnik B., Tepgec F., Karabey H. K., Uyguner Z. O.
JOURNAL OF ISTANBUL FACULTY OF MEDICINE-ISTANBUL TIP FAKULTESI DERGISI, vol.85, pp.162-169, 2022 (ESCI)

V. **Association between HBA locus copy number gains and pathogenic HBB gene variants**
Toksoy G., Akay N., Aghayev A., Karaman V., Avcı Ş., Kalaycı T., Altunoğlu U., Karakaş Z., Uyguner Z. O.
INTERNATIONAL JOURNAL OF MEDICAL BIOCHEMISTRY, vol.4, no.2, pp.91-95, 2021 (Peer-Reviewed Journal)

VI. **Association of HBA gene copy number gains with pathogenic HBB gene variants**
Toksoy G., Akay N., Aghayev A., Karaman V., Avcı Ş., Kalaycı T., Altunoğlu U., Karakaş Z., Uyguner Z. O.
International Journal of Medical Biochemistry, vol.4, no.2, pp.91-96, 2021 (Peer-Reviewed Journal)

VII. **MOLECULAR ANALYSIS OF FGFR1-3, TWIST1, MSX2, POR, FREM1 AND RAB23 GENES IN SYNDROMIC AND NON-SYNDROMIC CRANIOSYNOSTOSIS CASES**
Karaman V., Toksoy G., Karaman B., Kayserili Karabey H., Basaran S., Altunoglu U., Avci S., Uyguner Z. O.
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VIII. **SENDROMİK VE NON-SENDROMİK KRANİYOSİNOSTOZ OLGULARINDA FGFR1-3, TWIST1, MSX2, POR, FREM1 VE RAB23 GENLERİNİN MOLEKÜLER ANALİZİ**
Karaman V., TOKSOY G., KARAMAN B., KAYSERİLİ KARABEY H., BAŞARAN S., ALTUNOĞLU U., UYGUNER Z. O.
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Refereed Congress / Symposium Publications in Proceedings

I. **PIEZO1 İlişkili Dehidrate Herediter Stomasitoz-Herediter Kserositoz: Olgu Sunumu**
Konur E. N., Aslanger A. D., Ocak S., Karaman V., Uyguner Z. O., Yeşil Sayın G.
2. Ulusal HematoOnkoGenetik Kongresi , Gazimagusa, Cyprus (Kktc), 4 - 07 May 2023, pp.90

II. **Lenfoproliferatif Hastalıklarda Ayırıcı Tanıda Düşünülmesi Gereken Nadir Bir Sendrom: RAS İlişkili Otoimmün Lökoproliferatif Hastalık**
Yıldırım B. T., Akbaş S., Aslanger A. D., Karaman V., Yılmaz Y., Karaman S., Karaman B., Ünüvar A., Kılıç A., Uyguner Z. O.
2. Uluslararası Katılımlı Ulusal HematoOnkoGenetik Kongresi, Gazimagusa, Cyprus (Kktc), 4 - 07 May 2023, pp.111

III. **Osteogenezis Imperfekta Tanılı 15 Olgunun Moleküler Sonuçları**
Hacer Ö., Aslanger A. D., Kalaycı T., Güleç Ç., Demir K., Toksoy G., Karaman V., Öztürk A. P., Baş F., Yeşil Sayın G., et al.
15. Ulusal Tıbbi Genetik Kongresi, Muğla, Turkey, 09 November 2022, pp.149

IV. **Frank-Ter Haar Sendromu Tanili 3 Olgu ve Literatür Derlemesi**
Konur E. N., Aslanger A. D., Kalaycı T., Altunoğlu U., Karaman V., Yeşil Sayın G., Kayserili Karabey H., Uyguner Z. O.
15. Ulusal Tıbbi Genetik Kongresi, Muğla, Turkey, 09 November 2022, pp.91

V. **Gebelik Haftasına Göre Küçük Doğan (Sga) Çocuklarda Sebat Eden Boy Kısalığının Etiyolojisinin Genetik Analizler Ile Değerlendirilmesi**
Karaman V., Aslanger A. D., Konur E. N., Öztürk A. P., Toksoy G., Özsait Selçuk B. Ş., Baş F., Darendeliler F. F., Karaman B., Uyguner Z. O., et al.
15.ULUSLARARASI KATILIMLI, ULUSAL TIBBİ GENETİK KONGRESİ, Muğla, Turkey, 9 - 13 November 2022, pp.189

VI. **Investigation of Genes Associated with Multiple Pituitary Hormone Deficiencies via Next Generation Sequencing Technology**
ÖZTÜRK A. P., TOKSOY G., BAŞ F., YAVAŞ ABALI Z., Bagirova G., KARAMAN V., YILDIZ M., ASLANGER A., YEŞİL SAYIN G., POYRAZOĞLU Ş., et al.
60th Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), Roma, Italy, 15 September 2022

VII. **Çoğul Hipofiz Hormon Eksikliklerinde İlişkili Genlerin Yeni Nesil Dizileme Teknolojisi İle Araştırılması**
ÖZTÜRK A. P., TOKSOY G., BAŞ F., YAVAŞ ABALI Z., KARAMAN V., YILDIZ M., POYRAZOĞLU Ş., UYGUNER Z. O.,

DARENDELİLER F. F.
XXV. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, 06 October 2021

VIII. **MECP2 Spektrumundan Etkilenmiş 27 Olgunun Klinik ve Moleküler Bulguları**
Kalaycı T., Aslanger A. D., Altunoğlu U., Toksoy G., Konur E. N., Avcı Ş., Karaman V., Karaman B., Yeşil Sayın G., Kayserili Karabay H., et al.
14. TIBBİ GENETİK KONGRESİ, 20 - 22 December 2020, vol.31, no.4, pp.53

IX. **Alport sendromlu 15 olgunun klinik ve moleküler bulguları**
Aslanger A. D., Yürük Yıldırım Z. N., Toksoy G., Aksu B., Durmaz D., Göksu Çetinkaya A. P., Kalaycı T., Çam Delebe E. Ö., Karaman V., Yavuz S., et al.
14. TIBBİ GENETİK KONGRESİ, İstanbul, Turkey, 20 - 22 December 2020, vol.31, no.4, pp.49

X. **Ailevi Erken Puberte Olgularında MKRN3 ve DLK1 Genlerinin Dizilenmesi**
Karaman V., Karakılıç Özturan E., Baş F., Poyrazoğlu Ş., Başaran S., Darendeliler F. F., Uyguner Z. O.
XXIV. Ulusal Pediatrik Endokrinoloji ve Diyabet Çevrimiçi Kongresi , 30 October - 01 November 2020, pp.12

XI. **Analysis ofRUNX2mutations in four Turkish patients with Cleidocranial Dysplasia**
Mihci E., Guzel B. N., Toylu A., Karaman V., Aghayev A. R., Uyguner Z. O.
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XII. **Spectrum of Skeletal Abnormalities and Pathogenic RUNX2 Variants in 50 CleidocranialPatients from Turkey**
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XIII. **S-29 - Spectrum of Skeletal Abnormalities and Pathogenic RUNX2 Variants in 50 CleidocranialPatients from Turkey**
BERKAY E. G., ELKANOVA L., KALAYCI T., KARAMAN V., GÜNEŞ N., TOKSOY G., ALTUNOĞLU U., MIHÇI E., TAŞDELEN E., BAYRAMOĞLU Z., et al.
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XIV. **Novel FGFR2 variant in a Case with Crouzon Syndrome**
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XV. **Clinical features and genetic analyses of type III hereditary angioedema patients**
Gelincik A., Unal D., Demirturk M., Olgac M., Demir S., Toksoy G., Karaman V., Uyguner O., Colakoglu B., Buyukozturk S.
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XVI. **HBB gene mutation spectrum of beta-thalasemia patients from Turkey.**
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XVII. **Molecular Diagnostic Algorithm of Syndromic Craniosynostosis**
Karaman V., Toksoy G., Avcı Ş., Karaman B., Altunoğlu U., Başaran S., Kayserili Karabay H., Uyguner Z. O.
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XVIII. **Tip III Herediter Anjiödem Hastalarının Klinik ve Genetik Özelliklerinin Analizi**
GELİNCİK A., ÜNAL D., DEMİRTÜRK M., OLGAÇ M., DEMİR S., TOKSOY G., KARAMAN V., UYGUNER Z. O., ÇOLAKOĞLU B., BÜYÜKÖZTÜRK S.
XX. Ulusal Allerji ve Klinik immunoloji Kongresi, Antalya, Turkey, 2 - 06 November 2013, pp.51

XIX. **The first case diagnosed as both type III hereditary angioedema and familial Mediterranean fever**
Buyukozturk S., Gelincik a., Demirturk M., Unal D., Colakoglu B., Uyguner Z., Karaman V.
World Allergy and Asthma Congress of the European-Academy-of-Allergy-and-Clinical-Immunology and World-Allergy-Organization, Milan, Italy, 22 - 26 June 2013, vol.68, pp.250

XX. **Molecular Test Results of Syndromic Craniosynostosis Patients: genotype-phenotype correlations**
Karaman V., Altunoğlu U., Toksoy G., Karaman B., Kayserili Karabay H.

European Human Genetics Conference 2013, Paris, France, 8 - 11 June 2013, vol.21, no.1, pp.99

XXI. **Molecular Test Results of Syndromic Craniosynostosis Patients:genotype-phenotype correlations**
Karaman V., Altunoğlu U., Toksoy G., KARAMAN B., KAYSERİLİ H., UYGUNER Z. O.
European Human Genetic Congress, France, 1 - 04 June 2013, pp.99

XXII. **Describing the actual status of a Turkish bone marrow registry**
Diler A. S., Karaman V., Yesildag I., Yalman N.
22nd European Immunogenetics and Histocompatibility Conference, Toulouse, France, 2 - 05 April 2008, pp.377-378

XXIII. **HLAmatch - a programming library for allele and haplotype frequency based HLA matching**
Karaman V.
22nd European Immunogenetics and Histocompatibility Conference, Toulouse, France, 2 - 05 April 2008, pp.320-321

XXIV. **Increased HLA DRB1*11 with cGvHD and relation to engraftment in bone marrow transplantation**
Karaman V., Senturk-Ciftci H., Yalman N., Diler A. S.
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XXV. **Relatedness of mixed lymphocyte culture test with patient-donor parameters in hematopoietic stem cell transplantation**
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20th European Immunogenetics and Histocompatibility Conference, Oslo, Norway, 8 - 11 June 2006, pp.571

Supported Projects

UYGUNER Z. O., PARMAN F. Y., TÜYSÜZ B., KARA B., AKÇAYA N. H., YEŞİL SAYIN G., TOKSOY G., BAGIROVA G., ULUDAĞ ALKAYA D., ASLANGER A. D., et al., Project Supported by Higher Education Institutions, Dinamik mutasyon hastalıkları için moleküler genetik tanı kitlerinin geliştirilmesi, 2022 - Continues

DARENDELİLER F. F., BAŞ F., ASLANGER A. D., KARAMAN V., TOKSOY G., YEŞİL SAYIN G., KARAMAN B., ÖZTÜRK A. P., Project Supported by Higher Education Institutions, Büyümede Yakalama Yapamayan Gestasyon Yaşına Göre Düşük Doğum Ağırlıklı (SGA) Çocuklarda Boy Kısalığı Etiyolojisinin Araştırılması, 2022 - Continues

Scientific Refereeing

JOURNAL OF ISTANBUL FACULTY OF MEDICINE-ISTANBUL TIP FAKULTESI DERGISI, National Scientific Refreed Journal, January 2023

JOURNAL OF ISTANBUL FACULTY OF MEDICINE-ISTANBUL TIP FAKULTESI DERGISI, National Scientific Refreed Journal, December 2022

Metrics

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H-Index (Scopus): 4

Awards

Karakiliç Özturan E., Karaman V., Gelmez M. Y., Poyrazoğlu Ş., Baş F., Uyguner Z. O., Darendeliler F. F., A cause of familial central precocious puberty: A Novel variant in the DLK1 gene and low serum DLK1 levels, ESPE Travel Grant To Attend The 60Th Annual ESPE Meeting 2022,Rome , Espe And The Programme Organising Committee , September 2022
Karaman V., Karakılıç Özturan E., Baş F., Poyrazoğlu Ş., Başaran S., Darendeliler F. F., Uyguner Z. O., Sequence of MKRN3 and DLK1 genes in cases with familial central precocious puberty, ESPE Registration Grant To Attend The 59Th Annual Espe Meeting Online , Espe And The Programme Organising Committee , September 2021
Karaman V., Karakılıç Özturan E., Baş F., Başaran S., Uyguner Z. O., Two cases with central precocious puberty caused by paternally inherited novel variants in DLK1gene, European Human Genetics Conference 2020, June 2020