

Prof. TÜLAY GÜRAN

Personal Information

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International Researcher IDs

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Publons / Web Of Science ResearcherID: AGF-1845-2022

ScopusID: 13612528100

Yoksis Researcher ID: 183446

Education Information

Post Doctorate of Medicine, Marmara University, School of Medicine, Internal Medical Sciences, Turkey 2004 - 2009

Expertise In Medicine, Marmara University, School of Medicine, Internal Medical Sciences, Turkey 1999 - 2004

Undergraduate, Istanbul University, Cerrahpaşa Tıp Fakültesi, Cerrahpaşa Tıp Pr. (İngilizce), Turkey 1992 - 1999

Research Areas

Health Sciences

Academic and Administrative Experience

Faz Koordinatörü, Marmara University, School of Medicine, Internal Medical Sciences, 2020 - Continues

Mezuniyet Sonrası Eğitim Komisyonu Üyesi, Marmara University, School of Medicine, Internal Medical Sciences, 2020 - Continues

Courses

HEMATOPOIETIC SYSTEM AND RELATED DISORDERS SOLUK BENİZ MODÜLÜ, Undergraduate, 2021 - 2022

Jury Memberships

Appointment to Academic Staff-Assistant Professorship, Appointment Academic Staff, Marmara Üniversitesi, December, 2022

Doctoral Examination, Doctoral Examination, Marmara Üniversitesi, November, 2022

Expertise In Medicine, Expertise In Medicine, Marmara Üniversitesi, October, 2022

Appointment to Academic Staff-Assistant Professorship, Appointment Academic Staff, Sağlık Bilimleri Üniversitesi, September, 2022

Doctoral Examination, Doctoral Examination, Marmara Üniversitesi, September, 2022

Expertise In Medicine, Expertise In Medicine, Marmara Üniversitesi, August, 2022

Associate Professor Exam, Associate Professor Exam, Marmara University, July, 2022

Award, marmara student congress, Marmara Üniversitesi, May, 2022
Award, european society of pediatric endocrinology research unit board member, Marmara University, May, 2022
Appointment to Academic Staff-Professorship, Appointment Academic Staff, Altınbaş Üniversitesi, May, 2022
Expertise In Medicine, Expertise In Medicine, Marmara Üniversitesi, April, 2022
Associate Professor Exam, Associate Professor Exam, Marmara University, March, 2022
Expertise In Medicine, Expertise In Medicine, Marmara Üniversitesi, November, 2021
Appointment to Academic Staff-Professorship, Appointment Academic Staff, Koç Üniversitesi, August, 2021
Expertise In Medicine, Expertise In Medicine, Marmara Üniversitesi, July, 2021
Associate Professor Exam, Associate Professor Exam, Sağlık Bilimleri Üniversitesi, May, 2021
Expertise In Medicine, Expertise In Medicine, Marmara Üniversitesi, April, 2021
Appointment to Academic Staff-Assistant Professorship, Appointment Academic Staff, Sağlık Bilimleri Üniversitesi, March, 2021
Appointment to Academic Staff-Professorship, Appointment Academic Staff, Sağlık Bilimleri Üniversitesi, February, 2021
Appointment to Academic Staff-Professorship, Appointment Academic Staff, Sağlık Bilimleri Üniversitesi, February, 2021
Appointment to Academic Staff-Assistant Professorship, Appointment Academic Staff, İstanbul Aydın Üniversitesi, March, 2020
Associate Professor Exam, Associate Professor Exam, Marmara Üniversitesi, March, 2020
Appointment to Academic Staff-Professorship, Appointment Academic Staff, Altınbaş Üniversitesi, December, 2019
Associate Professor Exam, Associate Professor Exam, Marmara Üniversitesi, March, 2019
Appointment to Academic Staff-Professorship, Appointment Academic Staff, Koç Üniversitesi, November, 2018
Associate Professor Exam, Associate Professor Exam, Marmara Üniversitesi, December, 2017
Associate Professor Exam, Associate Professor Exam, Pamukkale Üniversitesi, October, 2017

Designed Courses And Trainings

Güran T., Çocuk Endokrinolojisi Ve Onkolojisi Kesişim Noktaları Ortak Sempozyumu, November 2022
Güran T., 26.Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi', October 2022
Güran T., I-DSD Training Workshop, July 2022
Güran T., Pediatrik Endokrinoloji İleri Kursu (PEİK), December 2019
Güran T., Pediatrik Endokrinolojiye Giriş Kursu, April 2019

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Lack of NAD(P)+ transhydrogenase activity in patients with primary adrenal insufficiency due to NNT variants**
Francisco A., Goler A. M. Y., Navarro C. D. C., Onder A., Yildiz M., Demirkol Y. K., Yilmaz B. K., Menevse T. S., GÜRAN T., Castilho R. F.
European Journal of Endocrinology, vol.190, no.2, pp.130-138, 2024 (SCI-Expanded)
- II. **Hormonal control during infancy and testicular adrenal rest tumor development in males with congenital adrenal hyperplasia: a retrospective multicenter cohort study**
Schröder M. A. M., Neacşu M., Adriaansen B. P. H., Sweep F. C. G. J., Ahmed S. F., Ali S. R., Bachega T. A. S. S., Baronio F., Birkebæk N. H., de Bruin C., et al.
European journal of endocrinology, vol.189, no.4, pp.460-468, 2023 (SCI-Expanded)
- III. **Challenges in the management of a 7 years old child with thyrotropin-secreting pituitary adenoma and the review of the literature**
KIRKGÖZ T., Abali S., Seker A., GÜRPINAR TOSUN B., ELTAN M., Helvacioğlu D., HALILOĞLU B., KAYGUSUZ S. B., Yavas Abali Z., SEVEN MENEVŞE T., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.96, no.5, pp.527-537, 2023 (SCI-Expanded)
- IV. **Pubertal and Gonadal Outcomes in 46,XY Individuals with Partial Androgen Insensitivity Syndrome**

Raised as Girls

Guaragna-Filho G., Guerra-Junior G., Tadokoro-Cuccaro R., Hughes I. A., Barros B. A., Hiort O., Balsamo A., GÜRAN T., Holterhus P. M., Hannema S., et al.

SEXUAL DEVELOPMENT, vol.17, no.1, pp.16-25, 2023 (SCI-Expanded)

- V. **Decline in the Age of Menarche in Istanbul Schoolgirls Over the Last 12 Years**
GÜRAN T., HELVACIOĞLU D., TOSUN B. G., ABALI Z. Y., Ahr F., Arslan Y. t., Molla G., Şahin B., Sayar M. E., Atay Z., et al.
Journal of Clinical Research in Pediatric Endocrinology, vol.15, no.2, pp.154-159, 2023 (SCI-Expanded)
- VI. **Introduction.**
Guran T., Flück C. E.
Hormone research in paediatrics, vol.96, no.2, pp.115, 2023 (SCI-Expanded)
- VII. **Molecular analysis of MKRN3 gene in Turkish girls with sporadic and familial idiopathic central**
KIRKGÖZ T., KAYGUSUZ S. B., ALAVANDA C., Helvacioglu D., Abali Z. Y., GÜRPINAR TOSUN B., ELTAN M., SEVEN MENEVŞE T., GÜRAN T., ARMAN A., et al.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.36, no.4, pp.401-408, 2023 (SCI-Expanded)
- VIII. **An Overlooked Manifestation of Hypercortisolism-Cerebral Cortical Atrophy and Challenges in Identifying the Etiology of Hypercortisolism: A Report of 2 Pediatric Cases**
Eviz E., Mutlu G. Y., Akcay A. A., Erbey F., GÜRAN T., Hatun S.
HORMONE RESEARCH IN PAEDIATRICS, 2023 (SCI-Expanded)
- IX. **Venous Thrombosis in a Pseudohypoparathyroidism Patient with a Novel GNAS Frameshift Mutation and Complete Resolution of Vascular Calcifications with Acetazolamide Treatment**
Menevse T. S., Iwasaki Y., Abali Z. Y., Tosun B. G., Helvacioglu D., DOĞRU Ö., BUĞDAYCI O., Cyr S. M., GÜRAN T., BEREKET A., et al.
Hormone Research in Paediatrics, 2023 (SCI-Expanded)
- X. **Evaluation of Cardiopulmonary Fitness and Cognitive Functions in Children with Obesity**
Ozturk C. C., Toprakoglu H. S., Oruc F., Menevse T. S., Akbolat F. I., GÜRAN T., BAHADIR A. T., KASIMAY Ö.
ACTA PHYSIOLOGICA, vol.237, pp.30, 2023 (SCI-Expanded)
- XI. **Rare forms of congenital adrenal hyperplasia**
GÜRPINAR TOSUN B., GÜRAN T.
Clinical Endocrinology, 2023 (SCI-Expanded)
- XII. **Genotype of congenital adrenal hyperplasia patients with testicular adrenal rest tumor**
AYCAN Z., Keskin M., Lafci N. G., Savas-Erdeve S., BAŞ F., POYRAZOĞLU Ş., Ozturk P., PARLAK M., Ercan O., GÜRAN T., et al.
EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.65, no.12, 2022 (SCI-Expanded)
- XIII. **A homozygous Y443C variant in the RNPC3 is associated with severe syndromic congenital hypopituitarism and diffuse brain atrophy**
Bezen D., Kutlu O., Mouilleron S., Rizzoti K., Dattani M., Guran T., Yesil G.
AMERICAN JOURNAL OF MEDICAL GENETICS PART A, vol.188, no.9, pp.2701-2706, 2022 (SCI-Expanded)
- XIV. **Glucagon response to hypoglycemia during extended oral glucose tolerance test in children with cystic fibrosis and comparing with healthy peers**
HALILOĞLU B., SEVEN MENEVŞE T., GÜRPINAR TOSUN B., GÜRAN T., DEMİRCİOĞLU S., Ispir T., GÖKDEMİR Y., ERDEM ERALP E., BEREKET A.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.199-200, 2022 (SCI-Expanded)
- XV. **Single Nucleotide Polymorphisms (SNPs) Profile as Predictive Markers of Lifestyle Modification Outcomes in Pediatric Obesity Treatment**
Gawlik A., Sobalska-Kwapis M., Antosz A., Strapagiel D., Seweryn M., Shmoish M., BEREKET A., Wasniewska M., KIRKGÖZ T., DEMİRCİOĞLU S., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.231-232, 2022 (SCI-Expanded)
- XVI. **A rare cause of monogenic obesity: Schaaf-Yang syndrome due to a novel MAGEL2 gene variant**
Abali Z. Y., Ates E. A., GÜRAN T., BEREKET A., DEMİRCİOĞLU S.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.230, 2022 (SCI-Expanded)
- XVII. **3 beta-Hydroxysteroid Dehydrogenase Deficiency, Rare in the Diagnosis of Congenital Adrenal**

Hyperplasia: A Case Report

Bulus A. D., Yasartekin Y., GÜRAN T.

HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.449, 2022 (SCI-Expanded)

- XVIII. Primary adrenal insufficiency in a patient with biallelic QRSL1 mutations**
Dursun F., Genc H. M., Yılmaz Göler A. M., Tas I., Eser M., Pehlivanoglu C., Yılmaz B., Güran T.
EUROPEAN JOURNAL OF ENDOCRINOLOGY, vol.187, no.3, 2022 (SCI-Expanded)
- XIX. Wide phenotypical spectrum with the same karyotype: Mixed gonadal dysgenesis**
Seven Menevşe T., Gürpınar Tosun B., Helvacıoglu D., Abali Z. Y., Kirmizibekmez H., Dursun F., Demircioğlu S., Bereket A., Güran T.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.566, 2022 (SCI-Expanded)
- XX. Diagnostic Features and Risk Factors for Childhood Thyroid Cancers**
ŞAHİN P., GÜRPINAR TOSUN B., YUMUŞAKHUYLU A. C., GÜRAN T., Helvacıoglu D., Abali Z. Y., HALILOĞLU B., OYSU Ç., BEREKET A., DEMİRCİOĞLU S.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.407, 2022 (SCI-Expanded)
- XXI. P1-206 Primary adrenal insufficiency in a patient with biallelic QRSL1 mutations**
Güran T.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.2, pp.125, 2022 (SCI-Expanded)
- XXII. Differences due to the variant type in the inheritance pattern of BMP15 gene-related primary ovarian insufficiency: a girl with a homozygous null BMP15 gene variant**
Abali Z. Y., Ates E. A., ELTAN M., GÜRPINAR TOSUN B., BEREKET A., GÜRAN T., DEMİRCİOĞLU S.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.377-378, 2022 (SCI-Expanded)
- XXIII. Circulating mRNA and miRNA Signatures as Predictive Markers of Lifestyle Modification Outcomes in Pediatric Obesity Treatment**
Gawlik A., Shmoish M., BEREKET A., Wasniewska M., Antosz A., KIRKGÖZ T., DEMİRCİOĞLU S., GÜRAN T., Aversa T., Corica D., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.213, 2022 (SCI-Expanded)
- XXIV. Change of menarcheal age in schoolgirls living in Istanbul over the last 12 years**
Güran T., Alir F., Arslan Y. T., Molla G., Sahin B., Sayar M. E., Atay Z., Helvacıoglu D., Gürpınar Tosun B., Haliloğlu B., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.364, 2022 (SCI-Expanded)
- XXV. An International Study of the Association between Local Health Care Resources and Acute Adrenal Insufficiency Events in Children with Congenital Adrenal Hyperplasia**
Tseretopoulou X., Ali S. R., Bryce J., Navoda A., Birkebaek N. H., Baronio F., Bonfig W., Claahsen-van der Grinten H. L., Cools M., Darendeliler F., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.401-402, 2022 (SCI-Expanded)
- XXVI. Breast ultrasonography: How useful in the diagnosis of precocious puberty?**
Helvacıoglu D., BIYIKLI E., BUĞDAYCI O., DEMİRCİOĞLU S., GÜRAN T., BEREKET A.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.46-47, 2022 (SCI-Expanded)
- XXVII. Gonadal morphology in 46,XY gonadal dysgenesis: I-DSD Registry-based study**
Tadokoro-Cuccaro R., Hughes I., Cools M., van de Vijver K., de Mendonca B. B., Domenice S., Batista R. L., Dallago R. T., Gomes N. L., Costa E. F., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.44-45, 2022 (SCI-Expanded)
- XXVIII. Urinary steroid metabolite ratios: sex- and age-dependent changes and use for the differential diagnosis of inborn steroidogenesis disorders**
Baranowski E. S., GÜRAN T., Gilligan L. C., Shaheen F., Utari A., Faradz S. M. H., Van Herwaarden A. E., Claahsen-van der Grinten H. L., Taylor A. E., Shackleton C. H. L., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.114, 2022 (SCI-Expanded)
- XXIX. A homozygous Y443C variant in the RNPC3 is associated with severe syndromic congenital hypopituitarism and diffuse brain atrophy**
Bezen D., Kutlu O., Mouilleron S., Rizzoti K., Dattani M., GÜRAN T., Yesil G.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.352-353, 2022 (SCI-Expanded)

- XXX. **Etiological analysis of hypophosphatemia: A single-center experience**
Eltan M., Alavanda C., Abali Z. Y., Bayramoglu E., Kaygusuz S. B., Helvacioğlu D., Tosun B. G., Menevse T. S., Ata P., Guran T., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.141-142, 2022 (SCI-Expanded)
- XXXI. **Basal cortisol measurements in the prediction of low-dose ACTH stimulation test outcomes**
Gacemer H., Gürpınar Tosun B., Abali Z. Y., Helvacioğlu D., Haliloğlu B., Demircioğlu S., Bereket A., Güran T.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.139, 2022 (SCI-Expanded)
- XXXII. **Gonadal morphology in 46,XY gonadal dysgenesis: I-DSD Registry-based study**
Güran T.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.Suppl 2, pp.44, 2022 (SCI-Expanded)
- XXXIII. **Predictors of surgical outcomes in boys with hypospadias**
Scougall K., Bryce J., Baronio F., Boal R. L., Castera R., Castro S., Cheetham T., Costa E. C., Darendeliler F., Davies J., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.376-377, 2022 (SCI-Expanded)
- XXXIV. **Low-dose ACTH Stimulation Test: Comparison of Cortisol Response at 30, 40, and 60 Minutes**
Gürpınar Tosun B., Arıkan H., Demircioğlu S., Bereket A., Güran T.
HORMONE RESEARCH IN PAEDIATRICS, vol.95, no.SUPPL 2, pp.117-118, 2022 (SCI-Expanded)
- XXXV. **A retrospective analysis of endocrine disease in sphingosine-1-phosphate lyase insufficiency: case series and literature review**
Maharaj A., Kwong R., Williams J., Smith C., Storr H., Krone R., Braslavsky D., Clemente M., Ram N., Banerjee I., et al.
ENDOCRINE CONNECTIONS, vol.11, no.8, 2022 (SCI-Expanded)
- XXXVI. **Adrenal steroids reference ranges in infancy determined by LC-MS/MS**
Enver E. O., Vatansever P., Guran O., Bilgin L., Boran P., Demircioğlu S., Haklar G., Bereket A., Güran T.
PEDIATRIC RESEARCH, vol.92, no.1, pp.265-274, 2022 (SCI-Expanded)
- XXXVII. **46,XY Partial gonadal dysgenesis; diagnosis and long-term outcome at puberty**
Cuccaro R. T., Hughes I., Cools M., van de Vijver K., de Mendonca B. B., Domenice S., Batista R. L., Dallago R. T., Gomes N. L., Costa E. F., et al.
SEXUAL DEVELOPMENT, vol.16, no.SUPPL 1, pp.33-35, 2022 (SCI-Expanded)
- XXXVIII. **Evaluation of thyroid function and metabolic parameters in obese and overweight children: A prospective case-control study.**
Kartal A. T., Bozaykut A., Sezer R. G., Güran T.
INVESTIGACION CLINICA, vol.63, no.2, pp.126-136, 2022 (SCI-Expanded)
- XXXIX. **Homozygosity for a novel INHA mutation in two male siblings with hypospadias, primary hypogonadism, and high normal testicular volume**
Guran T., Ates E. A., Eltan M., Sahin B., Tosun B. G., Seven Menevşe T., Geckinli B. B., Greenfield A., Turan S., Bereket A.
SEXUAL DEVELOPMENT, vol.16, no.SUPPL 1, pp.61-62, 2022 (SCI-Expanded)
- XL. **Treatment of congenital adrenal hyperplasia in children aged 0-3 years: a retrospective multicenter analysis of salt supplementation, glucocorticoid and mineralocorticoid medication, growth and blood pressure**
Neumann U., Van Der Linde A., Krone R. E., Krone N. P., Guven A., Güran T., Elsedfy H., Poyrazoglu S., Darendeliler F., Bachega T. A. S. S., et al.
EUROPEAN JOURNAL OF ENDOCRINOLOGY, vol.186, no.5, pp.587-596, 2022 (SCI-Expanded)
- XLI. **Familial Hypomagnesemia with Hypercalciuria and Nephrocalcinosis Due to CLDN16 Gene Mutations: Novel Findings in Two Cases with Diverse Clinical Features**
Eltan M., Abali Z. Y., Turkyilmaz A., Gökçe İ., Abali S., Alavanda C., Arman A., Kırkgöz T., Güran T., Hatun S., et al.
CALCIFIED TISSUE INTERNATIONAL, vol.110, no.4, pp.441-450, 2022 (SCI-Expanded)
- XLII. **Steroid hormone profiles and molecular diagnostic tools in pediatric patients with non-CAH primary adrenal insufficiency.**
Seven Menevşe T., Kendir Demirkol Y., Gurpınar Tosun B., Bayramoglu E., Yildiz M., Acar S., Erisen Karaca S., Orbak Z., Onder A., Sobu E., et al.

The Journal of clinical endocrinology and metabolism, vol.107, 2022 (SCI-Expanded)

- XLIII. **A novel deletion involving the first GNAS exon encoding Gs α causes PHP1A without methylation changes at exon A/B**
Campbell D., Reyes M., Kaygusuz S. B., Abalı S., Güran T., Bereket A., Kagami M., Turan S., Jüppner H.
Bone, vol.157, 2022 (SCI-Expanded)
- XLIV. **Homozygosity for a novel INHA mutation in two male siblings with hypospadias, primary hypogonadism, and high-normal testicular volume.**
Arslan Ateş E., Eltan M., Sahin B., Gurpinar Tosun B., Seven Menevse T., Geckinli B. B., Greenfield A., Turan S., Bereket A., Güran T.
European journal of endocrinology, vol.186, no.5, 2022 (SCI-Expanded)
- XLV. **Efficacy of the Novel Degludec/Aspart Insulin Co-formulation in Children and Adolescents with Type 1 Diabetes: A Real-life Experience with 1-year IDeg/Asp Therapy in Poorly Controlled and Non-compliant Patients.**
Kirkgoz T., Eltan M., Kaygusuz S. B., Yavas Abalı Z., Helvacioğlu D., Seven Menevse T., Gurpinar Tosun B., Guran T., Bereket A., Turan S.
Journal of clinical research in pediatric endocrinology, vol.14, pp.10-16, 2022 (SCI-Expanded)
- XLVI. **Systematic genetic testing for recessively inherited monogenic diabetes: a cross-sectional study in paediatric diabetes clinics**
Patel K. A., Ozbek M. N., Yildiz M., GÜRAN T., Kocyigit C., ACAR S., ŞIKLAR Z., Atar M., Colclough K., Houghton J., et al.
DIABETOLOGIA, vol.65, no.2, pp.336-342, 2022 (SCI-Expanded)
- XLVII. **Broad-spectrum XX and XY gonadal dysgenesis in patients with a homozygous L193S variant in PPP2R3C.**
Cicek D., Warr N., Yesil G., Kocak Eker H., Bas F., Poyrazoglu S., Darendeliler F., Direk G., Hatipoglu N., Eltan M., et al.
European journal of endocrinology, vol.186, pp.65-72, 2022 (SCI-Expanded)
- XLVIII. **Catch-up growth and discontinuation of fludrocortisone treatment in aldosterone synthase deficiency.**
Gurpinar Tosun B., Kendir Demirkol Y., Seven Menevse T., Kaygusuz S. B., Ozbek M. N., Altincik S. A., Mammadova J., Cayir A., Doger E., Bayramoglu E., et al.
The Journal of clinical endocrinology and metabolism, vol.107, 2022 (SCI-Expanded)
- XLIX. **Non-hormonal Clitoromegaly due to Clitoral Priapism Caused by Appendicitis/Appendectomy.**
Gurpinar Tosun B., Karagozlu Akgul A., Almus E., Abidoglu S., Turan S., Bereket A., Guran T.
Journal of clinical research in pediatric endocrinology, no.4, 2021 (SCI-Expanded)
- L. **Dysgenesis and Dysfunction of the Pancreas and Pituitary Due to FOXA2 Gene Defects.**
Kaygusuz S. B., Arslan Ates E., Vignola M. L., Volkan B., Geckinli B. B., Turan S., Bereket A., Gaston-Massuet C., Guran T.
The Journal of clinical endocrinology and metabolism, vol.106, no.10, 2021 (SCI-Expanded)
- LI. **Is quail egg a potential endocrine disruptor?**
Sürekli Karakuş Ö., Arabacı Tamer S., Levent H. N., Kaygusuz S. B., Demircioğlu S., Akakın D., Güran T., Yegen B., Bereket A.
HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.364, 2021 (SCI-Expanded)
- LII. **The exon 3 deleted/full length growth hormone receptor polymorphism and response to GH therapy in GH deficiency and Turner syndrome: a multicenter study**
Darendeliler F. F., Bas F., Bozkurt N., Uzunhan O., Aycan Z., Cetinkaya E., BERBEROĞLU M., ŞIKLAR Z., Ocal G., DARCAN Ş., et al.
HORMONE RESEARCH, vol.72, pp.109, 2009 (SCI-Expanded)
- LIII. **A novel missense mutation in the first extracellular loop of the neurokinin B receptor causes hypogonadotropic hypogonadism**
GÜRAN T., Tolhurst G., BEREKET A., Porter K., Turan S., Gribble F. M., KOTAN L. D., Akcay T., Atay Z., CANAN H., et al.
HORMONE RESEARCH, vol.72, pp.402, 2009 (SCI-Expanded)

Articles Published in Other Journals

- I. **Predictors of surgical complications in boys with hypospadias: Data from an international registry**
Scougall K., Bryce J., Baronio F., Boal R. L., Castera J. R., Castro S., Cheetham T., Costa E. C., Darendeliler F., Davies J. H., et al.
World Journal of Pediatric Surgery, vol.6, no.4, 2023 (ESCI)
- II. **[KS-015] İstanbul'da Yaşayan Kız Çocuklarında Son 12 Yılda Meydana Gelen Menarş Yaşı Değişiminin Belirlenmesi**
Güran T.
XXVI. ULUSAL PEDIATRİK ENDOKRİNOLOJİ XXVI. VE DİYABET KONGRESİ, vol.1, no.1, pp.10, 2022 (Conference Book)
- III. **Yenidoğanda Konjenital Adrenal Hiperplazi Taraması**
Güran T.
Türkiye Klinikleri Pediatri Dergisi, no.1, pp.32-35, 2022 (Peer-Reviewed Journal)
- IV. **Dysosteosclerosis: Clinical and Radiological Evolution Reflecting Genetic Heterogeneity**
DEMİRCİOĞLU S., Mumm S., ALAVANDA C., Kaygusuz B. S., GÜRPINAR TOSUN B., ARMAN A., Huskey M., GÜRAN T., Duan S., BEREKET A., et al.
JBMR PLUS, vol.6, no.8, 2022 (ESCI)
- V. **Insights From Long-term Follow-up of a Girl With Adrenal Insufficiency and Sphingosine-1-Phosphate Lyase Deficiency**
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LVII. Endokrin Söyleşiler-Yeni Gen Keşfinde Yolculuk

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- CIV. Ailevi glukokortikoid eksikliği tip 2: MR̈ap geninde yeni mutasyon p.K30del**

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CVI. Mutations in SGPL1, causing sphingosine-1-phosphate lyase deficiency, cause a novel form of primary adrenal insufficiency with steroid resistant nephrotic syndrome.

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CX. Precocious Puberty in Patients with Primary Adrenal Insufficiency due to Melanocortin Receptor 2 Mutation

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CXIII. b hCG from an Occult Source Causing Peripheral Precocious Puberty Identification of the Tumour 6 Years After Presentation

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CXIV. Mutations in SGPL1 cause a novel form of primary adrenal insufficiency with steroid resistant nephrotic syndrome

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- CXVIII. **Reconsideration of Mid Parental Height Calculation**
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- CXXV. **Zebra baliginda zCyp21a2 geninin karakterizasyonu**
Güran T., Zaucker A., Taylor A., Doultinos D., Griffin A., Mueller F., Krone N.
XIX. Ulusal Pediatrik Endokrinoloji, İstanbul, Turkey, 22 - 25 October 2015
- CXXVI. **Merkezi Yenidogan Tarama Programi ile Tani Almis Konjenital Hipotiroidili Vakalarimizin İzlemi**
Baş S., Abalı S., Atay Z., Gurbanov Z., Haliloglu B., Güran T., Turan S., Bereket A.
19. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Antalya, Turkey, 22 - 24 October 2015
- CXXVII. **Nonklasik Konjenital Adrenal Hiperplazi Hastalarının Genotip Ve Fenotip Özellikleri**
Abalı S., Akcan N., Toksoy G., Atay Z., Uyguner Z. O., Baş F., Güran T., Baş S., Kırmızıpekmez H., Poyrazoğlu Ş., et al.
19. PEDIATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Antalya, Turkey, 22 - 24 October 2015
- CXXVIII. **Otozomal Resesif Osteogenezis İmpperfekta Populasyonumuzdaki Sıklığı Ve Genetik Nedenleri**
ABALI S., ARMAN A., ATAY Z., BAŞ S., GÜRAN T., GÖRMEZ Z., DEMİRCİ H., BEREKET A., TURAN S.
19. PEDIATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Turkey, 22 - 24 October 2015
- CXXIX. **Periferal Puberte Prekokslu 129 Çocukta Etiyolojik Dağılım Ve Klinik Özellikler**
Atay Z., Yeşilkaya E., Savaş Ş., Turan S., Akın L., Eren E., Döğer E., Aycan Z., Yavaş Abalı Z., Akıncı A., et al.
19. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Antalya, Turkey, 22 - 24 October 2015
- CXXX. **5 Reduktaz tip 2 eksikliğine yol acan yeni mutasyonların moleküler karakterizasyonu**
Güran T., Shafqat N., Taylor A., Bujalska I., Ivison H., Wiebke A.
XIX. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, İstanbul, Turkey, 22 - 25 October 2015
- CXXXI. **5 Reduktaz tip 3 eksikliğinin insan steroid metabolizmasına etkisi**
Güran T., Bas F., Gokcay G., Hughes B. A., Kayserili Karabey H., Shackleton C. H., Arlt W.
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- CXXXII. **Boy Kısıklığı Olan Hastalarda Özellikler ve Etiyolojik Dağılım Bir Çocuk Endokrinoloji Kliniği Verileri**
Özcan S., Abalı S., Atay Z., Haliloğlu B., Baş S., Öztürk G., Akçay T., Güran T., Turan S., Bereket A.
19. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Antalya, Turkey, 22 - 24 October 2015
- CXXXIII. **İdrar steroid profili**
Güran T.
XIX. ULUSAL PEDIATRİK ENDOKRİNOLOJİ KONGRESİ, İstanbul, Turkey, 20 - 22 October 2015
- CXXXIV. **5 Reduktaz tip 2 eksikliği tanısında GC MS ile steroid olcumlerninde diagnostik oranların normatif datasinin olusturulması**
Güran T., Taylor A., Hughes B. A., Oneill D., Cedric S., Arlt W.
XIX. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, İstanbul, Turkey, 20 - 22 October 2015
- CXXXV. **Factors Effecting Response to Growth Hormone Treatment in Children with Turner Syndrome**
Baş S., Abalı S., Atay Z., Haliloğlu B., Gurbanov Z., Güran T., Bereket A., Turan S.
54th Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), Barcelona, Spain, 1 - 03 October 2015
- CXXXVI. **Hereditary Vitamin D Resistant Rickets Report of Four Cases with Successful Use of Intermittent Intravenous Calcium Via Peripheral Route**
Abalı S., Tamura M., Atay Z., İşgüven Ş. P., Güran T., Haliloğlu T., Baş S., Isojima T., Turan S., Kitanaka S., et al.
54th Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), Barcelona, Spain, 1 - 03 October 2015
- CXXXVII. **Evaluating First Year Response and Final Height to Growth Hormone Treatment in Growth Hormone Deficiency Based on Peak GH Levels on Testing**
Abalı S., Baş S., Akbarzade A., Atay Z., Haliloğlu B., Güran T., Turan S., Bereket A.
54th Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), Barcelona, Spain, 1 - 03 October 2015
- CXXXVIII. **Aetiological Spectrum and Clinical Characteristics of 129 Children with Gonadotropin Independent Precocious Puberty A Nationwide Cohort Study**
Atay Z., Yeşilkaya E., Erdeve Ş., Akin L., Eren E., Döğler E., Aycan Z., Abalı Z., Akıncı A., Şıklar Z., et al.
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- CXXXIX. **Primary Adrenal Insufficiency in Children without Congenital Adrenal Hyperplasia Molecular and Clinical Characterisation of a Nationwide Cohort**
Güran T., Buonocore F., Saka N., Özbek M. N., Aycan Z., Bereket A., Baş F., Darcan Ş., Bideci A., Turan S., et al.
54th Annual Meeting of the European Society for Paediatric Endocrinology (ESPE), Barcelona, Spain, 1 - 03 October 2015
- CXL. **A novel mutation cdel209 in the proopiomelanocortin gene in a child with early onset obesity**
CETINKAYA S., GÜRAN T., ERDAL K., Keskin M., Sagsak E., Savas Erdeve S., Buonocore F., AYCAN Z.
54th ESPE (European Society of Pediatric Endocrinology) meeting, Barselona, Spain, 29 September - 04 October 2015, vol.84, pp.1-622
- CXLI. **Identification and functional characterization of ESR2 a new disease gene for 46 XY disorders of sex development DSD**
Baetens D., Güran T., De Cauwer L., Looijenga L., De Bosscher K., Cools M., De Baere E.
5th I-DSD Symposium, Ghent, Belgium, 11 - 13 June 2015, vol.84
- CXLII. **Pubertal Development in Individuals with Partial Androgen Insensitivity Syndrome PAIS Assigned Female Sex of Rearing**
Guaragnafillo G., Junior G. G., Darendeliler F., Balsamo A., Holterhus P., Güran T., Ahmed S., Quigley C.
5th I-DSD Symposium, Ghent, Belgium, 11 - 13 June 2015, vol.84
- CXLIII. **ANTLEY BİXLER SENDROMLU BİR OLGUMUZ**
Turan S., Bas S., Akay Tayfun G., Abalı S., Atay Z., Qurbanov Z., Cam S., Huriye Nursel E., Güran T., Bereket A.
Cocuk Endokrinoloji Dernegi 7.Olgu Sunumlari Toplantisi, İzmir, Turkey, 18 - 20 May 2015
- CXLIV. **PROOPIOMELANOKORTİN POMC GENİNDE YENİ MUTASYONLU ERKEN BAŞLANGIÇLI BİR OBEZİTE OLGUSU**

Cetinkaya S., Aycan Z., Güran T., Kurnaz E., Keskin M., Sağsak E., Savaş Erdeve Ş., Achermann J.
Ulusal Çocuk Endokrinoloji Derneği 7. Olgu sunumlari toplantısı, İzmir, Turkey, 18 - 20 May 2015

CXLV. Urinary steroid Profiling

Güran T.

Disorders of Sex Development (DSD)- the roles of genes and the environment'PhD course, Soro, Denmark, 26 April - 27 October 2015

CXLVI. CHRONIC MUCOCUTANEOUS CANDIDIASIS, AUTOIMMUNE THYROIDITIS AND CEREBRAL MYCOTIC ANEURISM; STAT1 MUTATION

Kiykim A., Aydinler E., Ozen A. O., Baris S., Guran T., Hsu P. A., Barlan I.

100th J Project Meeting, Antalya, Turkey, 12 - 14 March 2014, vol.34, pp.745

CXLVII. Hipoglisemi: Kistik Fibrozise bağlı diyabet tanısında CGMS (sürekli glukoz izleme sistemi) ile OGTT nin karşılaştırılması

Haliloğlu B., Gökdemir Y., Atay Z., Abalı S., Güran T., Karakoç F., Ersu R., Karadağ B. T., Turan S., Bereket A.

TTD 17. yıllık kongresi, Antalya, Turkey, 2 - 06 April 2014

CXLVIII. Yenidoğanda hipogliseminin nadir bir nedeni: SUR 1 gen mutasyonuna bağlı hiperinsülinizm MEMİŞOĞLU A., GÜRAN T., İNAN E., DEMİREL B., KARATEKİN G.

19. Ulusal Neonatoloji Kongresi, Turkey, 17 - 20 April 2011

CXLIX. Identification of novel dentin matrix protein-1 (DMP1) mutations in two unrelated kindreds with autosomal recessive hypophosphatemia

Turan S., Bastepe M., Bereket A., Akçay T., Guran T., Makitie O., Juppner H.

29th Annual Meeting of the American-Society-for-Bone-and-Mineral-Research, Hawaii, United States Of America, 16 - 19 September 2007, vol.22

CL. Sağlıklı bebeklerde serum alkali fosfataz değerleri

Topçu B., Turan S., Gökçe İ., Akçay T., Güran T., Bereket A.

11. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Konya, Turkey, 14 - 17 September 2006, pp.118

Supported Projects

Güran T., Project Supported by Private Organizations in Other Countries, An Observational, Prospective Natural History Study of Early-Onset Extreme Obesity Due to Bi-Allelic Loss-of-Function Mutations in the POMC, PCSK1 or LEPR Genes. , 2020 - 2021

Güran T., Turan S., Project Supported by Private Organizations in Other Countries, A Phase 3 Randomized, Double-Blind, Placebo-Controlled, Parallel Group Study to Evaluate the Safety and Efficacy of Denosumab in Pediatric Subjects With Glucocorticoid-induced Osteoporosis, 2017 - 2021

Contractual Researches

Güran T., Diurnal, A Randomized, Double-Blind, Active-Controlled, Phase 3 Study of Chronocort Compared with Immediate-Release Hydrocortisone Replacement Therapy in Participants Aged 16 Years and Over with Congenital Adrenal Hyperplasia., 2022 - 2023

Güran T., Demircioğlu S., NovoNordisk, Büyüme hormonu eksikliği olan ve daha önce büyüme hormonu tedavisi almamış prepubertal çocuklarda, haftada bir uygulanan NNC0195-0092 tedavisi ile, günlük uygulanan büyüme hormonu tedavisinin (Norditropin® FlexPro®) etkililik ve güvenlik karşılaştırmasının araştırıldığı, randomize, çok uluslu, aktif kontrollü, (açık etiketli), doz bulma, (çift kör), paralel gruplu bir çalışma, 2022 - 2023

Demircioğlu S., Güran T., Buğdaycı O., Amgen, A Phase 3 Randomized, Double-blind, Placebo-controlled, Parallel-group Study to Evaluate the Safety and Efficacy of Denosumab in Pediatric Subjects with Glucocorticoid-induced Osteoporosis (GiOP), 2018 - 2021

Activities in Scientific Journals

HORMONE RESEARCH IN PAEDIATRICS, Assistant Editor/Section Editor, 2021 - Continues

Hormone Research In Paediatrics, Special Issue Editor, 2021 - Continues

FRONTIERS IN GENETICS, Assistant Editor/Section Editor, 2022 - 2022

FRONTIERS IN GENETICS, Assistant Editor/Section Editor, 2022 - 2022

Memberships / Tasks in Scientific Organizations

European Society of Pediatric Endocrinology-Winter school Steering Committee, Board Member, 2022 - Continues, England

Türk Çocuk Endokrinolojisi ve Diyabet Derneği Adrenal Çalışma Grubu, Vice President, 2021 - Continues, Turkey

Türk Çocuk Endokrinoloji ve Diyabet Derneği, Secretary General, 2020 - Continues, Turkey

I-DSD Steering Committee, Member, 2020 - Continues, United Kingdom

European society of pediatric endocrinology DSD working group , Chairman of the Scientific Committee, 2019 - Continues, England

European Society of Pediatric Endocrinology Science Committee Expert Panel, Member of Science Committee, 2018 - Continues, Germany

Scientific Refereeing

endo-2023, Conference Paper (Full Text), December 2022

PLOS ONE, Journal Indexed in SCI-E, December 2022

IMPE-2023, Conference Paper (Full Text), December 2022

Endocrine Connections, Journal Indexed in SCI-E, November 2022

JOURNAL OF STEROID BIOCHEMISTRY AND MOLECULAR BIOLOGY, Journal Indexed in SCI-E, November 2022

PLOS ONE, Journal Indexed in SCI-E, October 2022

INTERNATIONAL JOURNAL OF ENDOCRINOLOGY, Journal Indexed in SCI-E, October 2022

FRONTIERS IN GENETICS, Journal Indexed in SCI-E, September 2022

JCRPE JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, Journal Indexed in SCI-E, September 2022

JOURNAL OF ENDOCRINOLOGICAL INVESTIGATION, Journal Indexed in SCI-E, September 2022

European society of Pediatric Endocrinology meeting-2022, Conference Paper (Full Text), September 2022

CLINICAL ENDOCRINOLOGY, Journal Indexed in SCI-E, September 2022

HORMONE RESEARCH IN PAEDIATRICS, Journal Indexed in SCI-E, August 2022

HORMONE RESEARCH IN PAEDIATRICS, Journal Indexed in SCI-E, June 2022

JCRPE JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, Journal Indexed in SCI-E, May 2022

JOURNAL OF CLINICAL ENDOCRINOLOGY AND METABOLISM, Journal Indexed in SCI-E, May 2022

FRONTIERS IN GENETICS, Journal Indexed in SCI-E, May 2022

JOURNAL OF CLINICAL ENDOCRINOLOGY AND METABOLISM, Journal Indexed in SCI-E, April 2022

PLOS ONE, Journal Indexed in SCI-E, April 2022

upk-2022, Conference Paper (Full Text), April 2022

EXPERIMENTAL AND CLINICAL ENDOCRINOLOGY AND DIABETES, Journal Indexed in SCI-E, April 2022

REVIEWS IN ENDOCRINE AND METABOLIC DISORDERS, Journal Indexed in SCI-E, February 2022

JOURNAL OF PEDIATRIC ENDOCRINOLOGY AND METABOLISM, Journal Indexed in SCI-E, February 2022

FRONTIERS IN ENDOCRINOLOGY, Journal Indexed in SCI-E, January 2022

JOURNAL OF STEROID BIOCHEMISTRY AND MOLECULAR BIOLOGY, Journal Indexed in SCI-E, January 2022

JOURNAL OF PEDIATRICS, Journal Indexed in SCI-E, October 2021

JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, Journal Indexed in SCI-E, September 2021

EUROPEAN JOURNAL OF ENDOCRINOLOGY, Journal Indexed in SCI-E, July 2021

Other International Funding Programs, European Society of Pediatric Endocrinology Research Fellowship, United Kingdom, May 2021

Other International Funding Programs, European Society of Pediatric Endocrinology Research fellowship, United Kingdom, May 2021

JOURNAL OF CLINICAL ENDOCRINOLOGY & METABOLISM, Journal Indexed in SCI-E, March 2021

JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, Journal Indexed in SCI-E, February 2021

Scientific Consultations

büyüsün.org (<https://buyusun.org/uzman-gorusleri/hasta-buyume-hormonu-tedavisi-almayi-reddederse-ne-yapmali/>), Project Consultancy, Marmara University, School of Medicine, Internal Medical Sciences, Turkey, 2022 - Continues
Sağlık bakanlığı, Scientific Consultancy, Marmara University, School of Medicine, Internal Medical Sciences, Turkey, 2017 - Continues

marmara student congress, Project Consultancy, Marmara University, School of Medicine, Internal Medical Sciences, Turkey, 2022 - 2022

Tasks In Event Organizations

Demircioğlu S., GÜRAN T., XXVI. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Scientific Congress, Turkey, Ekim 2022

Güran T., VIII. PEDIATRİK ENDOKRİNOLOJİYE GİRİŞ KURSU, Scientific Congress, Antalya, Turkey, Eylül 2022

Güran T., Steroids, Mass Spectrometry and Endocrinology – Past, Present and Future, Scientific Congress, Birmingham, England, Nisan 2022

Güran T., VII. PEDIATRİK ENDOKRİNOLOJİ İLERİ KURSU, Scientific Congress, İstanbul, Turkey, Şubat 2022

Güran T., 9. MARMARA PEDIATRİ KONGRESİ, Scientific Congress, İstanbul, Turkey, Şubat 2022

Metrics

Publication: 437

Citation (WoS): 2708

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H-Index (WoS): 26

H-Index (Scopus): 26

Congress and Symposium Activities

NovoCME Continuous Medical Education International Academy online, Moderator, İstanbul, Turkey, 2022

ÇEDD Uzaktan Eğitim Seminerleri, Moderator, İstanbul, Turkey, 2022

XXVI.ULUSAL PEDIATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Invited Speaker, Antalya, Turkey, 2022

XXVI. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Moderator, Antalya, Turkey, 2022

2. Cerrahpaşa Pediatri Günleri, Session Moderator, İstanbul, Turkey, 2022

European Society for Paediatric Endocrinology 2022 meeting, Moderator, Rome, Italy, 2022

The European Society for Paediatric Endocrinology, Session Moderator, Rome, Italy, 2022

The European Society for Paediatric Endocrinology (ESPE) meeting 2022, Session Moderator, Rome, Italy, 2022

NovoCME Continuous Medical Education International Academy online, Moderator, İstanbul, Turkey, 2022

ÇOCUK ENDOKRİNOLOJİSİ OLGU SUNUMLARI -11, Session Moderator, İstanbul, Turkey, 2022

18. uLUDAG PEDIATRİ KİS KONGRESİ, Session Moderator, Bursa, Turkey, 2022

9. Marmara Pediatri Kongresi, Session Moderator, İstanbul, Turkey, 2022

NovoCME Continuous Medical Education International Academy online, Moderator, İstanbul, Turkey, 2021

XXV. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Session Moderator, Antalya, Turkey, 2021

European Society for Paediatric Endocrinology 2021 meeting, Moderator, Liverpool, United Kingdom, 2021
8th I-DSD Symposium 2021, Session Moderator, Glasgow, United Kingdom, 2021
10.Olgu Sunumları Sempozyumu , Session Moderator, İzmir, Turkey, 2021

Invited Talks

Endokrin Hastalıkların Genetiği Seminerleri- 46,XY Primer Gonadal Yetmezlikte yeni bir gen: INHA, Seminar, ÇOCUK ENDOKRİNOLOJİSİ VE DİYABET DERNEĞİ, Turkey, October 2022
Ulusal yenidoğan KAH taraması, Conference, 66. Türkiye Milli Pediatri Kongresi, Turkey, October 2022
Cinsiyet Gelişim Bozukluğu Olgularında Endokrin Testlerin Yorumlanması - Tülay Güran
(<https://www.cocukendokrindiabetes.org/eogrenmeliste>), Seminar, ÇOCUK ENDOKRİNOLOJİSİ VE DİYABET DERNEĞİ, Turkey, September 2022
yearbook, Conference, The European Society for Paediatric Endocrinology, Italy, September 2022
Interpretation of endocrine tests , Conference, 9th I-DSD Symposium 2022, Switzerland, July 2022
ESPE DSD working group, Conference, 9th I-DSD Symposium 2022, Turkey, July 2022
Ulusal KAH taraması, Conference, 57. Türk Pediatri Kongresi - Kongre Uzmanı, Turkey, May 2022
Steroids, Mass Spectrometry and Endocrinology – Past, Present and Future, Conference, University of Birmingham, England, April 2022
Ulusal yenidoğan KAH taraması, Conference, Bursa Uludağ Üniversitesi, Turkey, March 2022
KAH Olgu Çalışması, Seminar, VII. Pediatrik Endokrinoloji İleri Kursu (PEİK), Turkey, February 2022
Yenidoğan KAH Taraması Toplantısı: Sorunlar - Çözüm Önerileri - Tülay Güran(<https://www.cocukendokrindiabetes.org/eogrenmeliste>), Seminar, ÇOCUK ENDOKRİNOLOJİSİ VE DİYABET DERNEĞİ, Turkey, January 2022
Türkiye`de Konjenital Adrenal Hiperplazi Yenidoğan Taraması: Verilerin Değerlendirilmesi, Conference, 65. Türkiye Milli Pediatri Kongresi, Turkey, November 2021
Optimizing Medical Therapy in Different Forms of Congenital Adrenal Hyperplasia , Seminar, KONIKA XVIII Medan 2021, Indonesia, October 2021
Common and rare adrenal causes of DSD, Seminar, BSPED DSD SIG Programme , United Kingdom, October 2021
46,XX cinsiyet gelişim bozukluklarına yaklaşım, Seminar, XXV. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, October 2021
Steroidogeneizde Aberan Androjen Üretimi ve Klinik Yansımaları, Seminar, XXV. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, October 2021
KONJENİTAL ADRENAL HİPERPLAZİNİN YÖNETİMİ , Seminar, VI. Pediatrik Endokrinoloji İleri Kursu, Turkey, June 2021
Optimizing Medical Therapy in Different Forms of Congenital Adrenal Hyperplasia , Seminar, Indian Society for Pediatric and Adolescent Endocrinology (ISPAE)Academics and Clinical Education Series (ISPAE - ACES), India, May 2021
Cocuklarda gecikmis ergenlik, Seminar, Endokrinoloji Talim, Turkey, April 2021
Steroidogeneizde Arka Yolak Hastalıklarının Klinik Yansımaları, Seminar, 4. Ege Endokrin Hastalıklar ve Genetik Sempozyumu, Turkey, March 2021
Konjenital Adrenal Hiperplazi Tanısında Zorluklar , Seminar, Endokrin Hastalıkların Genetiği Seminerleri, Turkey, January 2021
46, XX Cinsiyet gelişim bozukluklarına yaklaşım , Seminar, VI. PEDİATRİK ENDOKRİNOLOJİYE GİRİŞ KURSU , Turkey, November 2020
Geç tanı alan Turner sendromu olgusunda büyüme hormonu tedavisi ve puberte induksiyonu, Seminar, NordiPEVA Büyüyen Tecrübe Toplantısı, Turkey, October 2020
Adrenal Yetmezlik ve Yönetimi , Seminar, Çocuk Endokrinolojisinde Güncellemeler, Turkey, September 2020
Endokrin Söyleşiler-Yeni Gen Keşfinde Yolculuk, Seminar, 24. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, April 2020

Scholarships

Desteklenen Araştırma Ödülleri (Yurtiçi), Other Government Agencies, 2022 - 2023
BAPKO TTU-2022-10603, University, 2022 - 2022
Desteklenen Araştırma Ödülleri (Yurtiçi), Other Government Agencies, 2022 - 2022
Registration Grant, Other International Organizations, 2021 - 2021
Registration grant, Other International Organizations, 2021 - 2021
Registration grant, Other International Organizations, 2021 - 2021

Awards

Güran T., RESEARCH PARTICIPATION AWARD, 9Th I-Dsd Symposium 2022, July 2022
Güran T., Seven Menevse T., Turan S., Bereket A., Gulpinar Tosun B., Sozel sunum ucunculuk odulu , Turk Cocuk Endokrinoloji Ve Diyabet Dernegi , October 2021
Güran T., Darendeliler F., Yilin en iyi yayın odulu, Turk Cocuk Endokrinoloji Ve Diyabet Dernegi, October 2021
Yavas Abali Z., Güran T., Turan S., Bereket A., Eltan M., People`s choice poster award, European Society Of Pediatric Endocrinology, September 2021
Güran T., Turan S., Bereket A., Surekli Karakus O., Yegen B., Dr Tolga Koroglu özel odulu , 8. Marmara Pediatri Kongresi, February 2021
Güran T., Demircioğlu S., Bereket A., Eltan M., Gürpınar Tosun B., Kısa Sözel Bildiri Üçüncülük Ödülü- Biallelik PPP2R3C mutasyonları 46, XX ve 46, XY gonadal disgeneziye yol açar, Xxiv. Ulusal Pediatrik Endokrinoloji Ve Diyabet Çevrim İçi Kongresi, November 2020
Güran T., Demircioğlu S., Bereket A., Haklar G., Sözlü Bildiri Birincilik Ödülü Steroid 11 β -hidroksilaz eksikliği olan 100 çocuk hastanın klinik bulgularının genetik ve adrenokortikal hormon profili ile ilişkisinin değerlendirilmesi, Xxiv. Ulusal Pediatrik Endokrinoloji Ve Diyabet Çevrim İçi Kongresi, November 2020
Güran T., Demircioğlu S., Bereket A., Kaygusuz S. B., Eltan M., Tutar E., Gürpınar Tosun B., Seven Menevşe T., Poster Bildiri Birincilik Ödülü-Parsiyel Pankreatik Agenezi ve Sendromik Hipopituitarizm, Xxiv. Ulusal Pediatrik Endokrinoloji Ve Diyabet Çevrim İçi Kongresi, November 2020
Haliloğlu B., Hysenaj G., Atay Z., Güran T., Abalı S., Demircioğlu S., Ellard S., Bereket A., Best second oral presentation, Ulusal Pediatrik Endokrinoloji Kongresi Bilimsel Kurul, October 2013

Representation and Promotion Activities

Institutional Representation, Saglik Bakanligi, Turkey, Ankara, 2017 - 2022

Non Academic Experience

University of Birmingham Center for Endocrinology Diabetes and Metabolism
University of Cambridge Institute of Metabolic Sciences