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Foreign Languages

English, C1 Advanced

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Academic Titles / Tasks

Professor, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 1996 - Continues

Associate Professor, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 1989 - 1996

Assistant Professor, Istanbul University, Istanbul Medical Faculty, Division Of Medical Sciences , 1989 - 1989

Academic and Administrative Experience

İstanbul Üniversitesi, Büyüme-Gelişme Ve Çocuk Endokrinoloji Bilim Dalı, Büyüme-Gelişme Ve Çocuk Endokrinoloji Bilim Dalı, 2013 - Continues

İstanbul Üniversitesi, Çocuk Sağlığı Ve Hastalıkları Anabilim Dalı , Büyüme-Gelişme Ve Çocuk Endokrinoloji Bilim Dalı, 2012 - Continues

İstanbul Üniversitesi, Istanbul Medical Faculty, Büyüme-Gelişme Ve Çocuk Endokrinoloji Bilim Dalı, 2011 - Continues

Advising Theses

DARENDELİLER F. F. , İdyopatik boy kısalığında ALS düzeyleri, Expertise In Medicine, D.Tekcan(Student), 2005

DARENDELİLER F. F. , SGAlı prepubertal çocuklarda insülin direncinin araştırılması, Expertise In Medicine, Ö.Sancaklı(Student), 2003

DARENDELİLER F. F. , Büyüme hormonu eksikliğinde cins steroid uyarılı bazal ve uyarılmış IGF1 ve IGFBP3 düzeylerin değerlendirilmesi, Expertise In Medicine, C.Öcal(Student), 2002

DARENDELİLER F. F. , Büyüme hormonu eksikliğinde büyüme hormonu testlerinin tekrar değerlendirilmesi, Expertise In Medicine, İ.Spinu(Student), 2000

DARENDELİLER F. F. , Konjenital hipotiroidide büyümeyi etkileyen çeşitli parametreler ve büyümenin izlenmesi, Expertise In Medicine, M.Yıldırım(Student), 1994

DARENDELİLER F. F. , Malign hematolojik hastalıklarda gonadal işlevler, Expertise In Medicine, Ö.Özdemir(Student), 1994

DARENDELİLER F. F. , 1 yaşındaki çocuklarda ebeveyn boyuna göre erişkin boyunun tahmini, Expertise In Medicine, B.Bayraktar(Student), 1992

Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Comparison of National Growth Standards for Turkish Infants and Children with World Health Organization Growth Standards**
Bundak R., Abali Z. Y. , Furman A., Darendeliler F., Gokcay G., Bas F., Gunoz H., Neyzi O.
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- II. **The Impact of the CEDD-NET on the Evaluation of Rare Disorders: A Multicenter Scientific Research Platform in the Field of Pediatric Endocrinology**
ÖZEN S., Ata A., Darendeliler F.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.14, no.2, pp.216-220, 2022 (Journal Indexed in SCI)
- III. **Mutations in AR or SRD5A2 Genes: Clinical Findings, Endocrine Pitfalls, and Genetic Features of Children with 46,XY DSD**
Akcan N., Uyguner O., Bas F., Altunoglu U., Toksoy G., Karaman B., Avci S., Abali Z. Y. , Poyrazoglu S., Aghayev A., et al.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.14, no.2, pp.153-171, 2022 (Journal Indexed in SCI)
- IV. **Impact of Smoking, Obesity and Maternal Diabetes on SHBG Levels in Newborns.**
Aydin B. K. , Yasa B., Moore J. P. , Yasa C., Poyrazoglu Ş., Bas F., Coban A., Darendeliler F. F. , Winters S. J.
Experimental and clinical endocrinology & diabetes : official journal, German Society of Endocrinology [and] German Diabetes Association, vol.130, no.5, pp.335-342, 2022 (Peer-Reviewed Journal)
- V. **Pelvic and breast ultrasound abnormalities and associated metabolic disturbances in girls with premature pubarche due to adrenarche.**
Aydin B. K. , Kadioglu A., Kaya G. A. , Devcioglu E., Baş F., Poyrazoğlu Ş., Gökçay E. G. , Darendeliler F. F.
Clinical endocrinology, vol.96, no.3, pp.339-345, 2022 (Peer-Reviewed Journal)
- VI. **Growth and Pubertal Features in a Cohort of 83 Patients with Osteogenesis Imperfecta.**
Ozturk A. P. , Dudakli A., Ozturan E., Poyrazoglu Ş., Bas F., Darendeliler F.
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- VII. **Growth, puberty and testicular function in boys born small for gestational age with a nonspecific disorder of sex development.**
Tack L. J. W. , van der Straaten S., Riedl S., Springer A., Holterhus P., Hornig N. C. , Kolesinska Z., Niedziela M., Baronio F., Balsamo A., et al.
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- VIII. **Long-term Follow-up of a Toddler with Papillary Thyroid Carcinoma: A Case Report with a Literature Review of Patients Under 5 Years of Age.**
Öztürk A. P. , Karakılıç Özturan E., Gün Soysal F., Ünal S., Işık G., Yegen G., Önder S., Yıldız M., Poyrazoğlu Ş., Baş F., et al.
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- IX. **Recommendations for Clinical Decision-making in Children with Type 1 Diabetes and Celiac Disease: Type 1 Diabetes and Celiac Disease Joint Working Group Report.**
Hatun Ş., Dalgıç B., Gökşen D., Aydoğdu S., Savaş Erdeve Ş., Kuloğu Z., Doğan Y., Aycan Z., Yeşiltepe Mutlu G., Uslu Kızılkın N., et al.
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- X. **Broad-spectrum XX and XY gonadal dysgenesis in patients with a homozygous L193S variant in PPP2R3C.**

ÇİÇEK D., Warr N., Yesil G., Eker H. K. , Bas F., Poyrazoglu Ş., Darendeliler F., DİREK G., HATİPOĞLU N., ELTAN M., et al.

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- XI. **Evaluation of the Efficacy and Safety of 3 Different Management Protocols in Pediatric Diabetic Ketoacidosis.**
Akcan N., Uysalol M., Kandemir I., Soydemir D., Abali Z., Poyrazoglu Ş., Bas F., Bundak R., Darendeliler F. F.
Pediatric emergency care, vol.37, 2021 (Peer-Reviewed Journal)
- XII. **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.
HORMONE RESEARCH IN PAEDIATRICALS, vol.94, no.SUPPL 1, pp.167-168, 2021 (Peer-Reviewed Journal)
- XIII. **Evaluation of Persistent Short Stature in Children Born Small for Gestational Age without Catch-up Growth**
Ozturk A. P. , Ozturan K., Poyrazoglu S., Bas F., Darendeliler F.
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- XIV. **Long-Term Cardiometabolic Morbidity in Young Adults with Classic 21-Hydroxylase Deficiency Congenital Adrenal Hyperplasia**
Righi B., Ali S. R. , Bryce J., Tomlinson J. W. , Bonfig W., Baronio F., Costa E. C. , Guaragna Filho G., T'Sjoen G., Cools M., et al.
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- XV. **Growth and Puberty in Patients with Osteogenesis Imperfecta**
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- XVI. **Response to growth hormone therapy with high IGF-1-levels and severe insulin resistance in two-cases with SOFT syndrome: A novel homozygous mutation in POC1A**
Karakilic-Ozturan E., Altuoglu U., Ozturk A. P. , Toksoy G., Turgut G. T. , Poyrazoglu S., Bas F., Uyguner O., Darendeliler F.
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- XVII. **Biallelic PPP2R3C mutations are associated with partial and complete gonadal dysgenesis in 46,XY and 46,XX individuals**
ÇİÇEK D., Warr N., Yesil G., Eker H. K. , Bas F., Poyrazoglu S., Darendeliler F., DİREK G., HATİPOĞLU N., ELTAN M., et al.
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- XVIII. **Clinical and hormonal profiles correlate with molecular characteristics in patients with 11β-hydroxylase deficiency.**
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- XIX. **Broad range of phenotypes in an international cohort of 75 DSD individuals with SF-1/NR5A1 variants**
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- XX. **Evaluation of early puberty in boys and girls with Silver-Russell Syndrome: Discordance between testicular growth and pituitary-gonadal hormones in male cases**
Yildiz M., Bas F., Karaman B., Poyrazoglu S., Basaran S., Darendeliler F.
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- XXI. **Phenotypic Characteristics of Patients with 45,X/46,XY Mosaicism: Growth, Gonadal Pathology and Tumour Risk**
Poyrazoglu S., Bas F., Karaman B., Yildiz M., Basaran S., Darendeliler F.
HORMONE RESEARCH IN PAEDIATRICALS, vol.94, no.SUPPL 1, pp.86-87, 2021 (Peer-Reviewed Journal)
- XXII. **Effects of growth hormone therapy on serum concentrations of IGF-1 in patients with Turner syndrome: High IGF-1 concentrations despite optimal dose?**

- Ozturan E. K. , Karagoz N., Ceylaner S., Ozturk A. P. , Al A. D. K. , Abali Z. Y. , Poyrazoglu S., Bas F., Darendeliler F. HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.317-318, 2021 (Peer-Reviewed Journal)
- XXIII. **Cranial MRI abnormalities and long-term follow-up of the lesions in 770 girls with Central Precocious Puberty.**
Helvacioğlu D., Demircioğlu Turan S., Güran T., Atay Z., Dağcınar A., Bezen D., Karakılıç Özturan E., Darendeliler F. F. , Yüksel A., Dursun F., et al.
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- XXIV. **Monogenic Childhood Diabetes: Dissecting Clinical Heterogeneity by Next-Generation Sequencing in Maturity-Onset Diabetes of the Young.**
Kanca Demirci D., Darendeliler F. F. , Poyrazoglu Ş., Al A. D. K. , Gül N., Tutuncu Y., Gulfidan G., Arğa K. Y. , Cacina C., Öztürk O., et al.
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- XXV. **Clinical Characteristics of 46,XX Males with Congenital Adrenal Hyperplasia**
Savaş-Erdeve Ş., Aycan Z., Çetinkaya S., Ozturk A. P. , Bas F., Poyrazoglu Ş., Darendeliler F. F. , Ozsu E., Sıklar Z., Demiral M., et al.
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- XXVI. **LRBA deficiency: a rare cause of type 1 diabetes, colitis, and severe immunodeficiency.**
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- XXVII. **Gonadectomy in conditions affecting sex development: a registry-based cohort study.**
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- XXVIII. **International practice of corticosteroid replacement therapy in congenital adrenal hyperplasia: data from the I-CAH registry.**
Bacila I., Freeman N., Daniel E., Sandrk M., Bryce J., Ali S. R. , Yavas Abali Z., Atapattu N., Bachega T. A. , Balsamo A., et al.
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- XXIX. **Growth and relationship of phenotypic characteristics with gonadal pathology and tumour risk in patients with 45, X/46, XY mosaicism.**
Poyrazoglu Ş., Bas F., Karaman B., Yildiz M., Başaran S., Darendeliler F. F.
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- XXX. **HEART AND AORTA ANOMALIES IN TURNER SYNDROME AND RELATION WITH KARYOTYPE**
Al Kardelen A. D. , Gencay G., BAYRAMOĞLU Z., Aliyev B., Karakilic-Ozturan E., POYRAZOĞLU Ş., NİŞLİ K., BAŞ F., DARENDELİLER F. F.
ACTA ENDOCRINOLOGICA-BUCHAREST, vol.17, no.1, pp.124-130, 2021 (Journal Indexed in SCI)
- XXXI. **Testosterone Therapy and Its Monitoring in Adolescent Boys with Hypogonadism: Results of an International Survey from the I-DSD Registry.**
Stancampiano M. R. , Lucas-Herald A. K. , Bryce J., Russo G., Barera G., Balsamo A., Baronio F., Bertelloni S., Valiani M., Cools M., et al.
Sexual development : genetics, molecular biology, evolution, endocrinology, embryology, and pathology of sex determination and differentiation, vol.15, no.4, pp.236-243, 2021 (Peer-Reviewed Journal)
- XXXII. **Real World Estimates Of Adrenal Insufficiency Related Adverse Events In Children With Congenital Adrenal Hyperplasia.**
Ali S., Bryce J., Haghpanahan H., Lewsey J., Tan L., Atapattu N., Birkebaek N., Blankenstein O., Neumann U., Balsamo A., et al.
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- XXXIII. **Loss-of-function variants in SEMA3F and PLXNA3 encoding semaphorin-3F and its receptor plexin-A3 respectively cause idiopathic hypogonadotropic hypogonadism**
KOTAN L. D. , Ternier G., Cakir A. D. , EMEKSİZ H. C. , TURAN İ., Delpouve G., Kardelen A. D. , Ozcabi B., Isik E., Mengen E., et al.

GENETICS IN MEDICINE, 2021 (Peer-Reviewed Journal)

- XXXIV. **Care and Support of Children with Type 1 Diabetes at School: The Turkey Experience.**
Hatun Ş., Yeşiltepe Mutlu G., Gökçe T., Avcı Ö., Yardım N., Aycan Z., Darendeliler F. F.
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- XXXV. **Factors affecting physicians' perception of the overuse of antibiotics.**
Camcioglu Y., Sener O., Aksaray N., Darendeliler F. F. , Hasanoglu E.
Medecine et maladies infectieuses, vol.50, pp.652-657, 2020 (Peer-Reviewed Journal)
- XXXVI. **An evaluation of the knowledge and attitudes of school staff related to diabetes care at school: The 10th year of the "diabetes program at school" in Turkey**
Gokce T., Sakarya S., Muradoglu S., Mutlu G. Y. , Can E., Cemhan K., Kurtulmus M. F. , Gulsen M., Aycan Z., Darendeliler F., et al.
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- XXXVII. **Neonatal Screening for Congenital Adrenal Hyperplasia in Turkey: Outcomes of Extended Pilot Study in 241,083 Infants**
GÜRAN T., Tezel B., Cakir M., AKINCI A., ORBAK Z., Keskin M., Eklioglu B. S. , Ozon A., Ozbek M. N. , KARAGÜZEL G., et al.
Journal of clinical research in pediatric endocrinology, vol.12, no.3, pp.287-294, 2020 (Journal Indexed in SCI Expanded)
- XXXVIII. **Pituitary Iron Deposition and Endocrine Complications in Patients with β -Thalassemia: From Childhood to Adulthood.**
Karadag S., Karakas Z., Yilmaz Y., Gul N., Demir A., Bayramoglu Z., Darendeliler F. F. , Dursun M.
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- XXXIX. **New Features for Child Metrics: Further Growth References and Blood Pressure Calculations**
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- XL. **Superb Microvascular Imaging in the Evaluation of Pediatric Graves Disease and Hashimoto Thyroiditis.**
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- XLI. **What is the evidence for beneficial effects of growth hormone treatment beyond height in short children born small for gestational age? A review of published literature.**
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- XLII. **Plasma Renin Measurements are Unrelated to Mineralocorticoid Replacement Dose in Patients With Primary Adrenal Insufficiency**
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- XLIII. **A Novel Homozygous Mutation of the Acid-Labile Subunit (IGFALS) Gene in a Male Adolescent**
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- XLIV. **Response to Letter to the Editor: "Clinical but Not Histological Outcomes in Males With 45,X/46,XY Mosaicism Vary Depending on Reason for Diagnosis".**
Ljubicic M. L. , Jorgensen A., Ribeiro de Andrade J. G. , Balsamo A., Bertelloni S., Cools M., Cuccaro R. T. , Darendeliler F. F. , Fluck C. E. , Grinspon R. P. , et al.
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- XLV. **Clinical but Not Histological Outcomes in Males With 45,X/46,XY Mosaicism Vary Depending on Reason for Diagnosis.**
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- XLVI. **Editorial: Hot Topics of Debate on Turner Syndrome: Growth, Puberty, Cardiovascular Risks, Fertility and Psychosocial Development.**
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- XLVII. **Genotype-Phenotype Correlation and Clinical Findings in 145 Patients with Congenital Adrenal Hyperplasia: Single Centre Experience**
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- XLVIII. **Genetic Evaluation of Idiopathic Short Stature**
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- XLIX. **Serum endocan levels as a marker of endothelial dysfunction in Turner syndrome and correlation with cardiac findings**
Gencay A. G. , Darendeliler F. F. , Nisli K., Karaca S., Kardelen A. D. , Poyrazoglu S., Bas F.
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- L. **Influence of salt supplementation on drug therapy in children with congenital adrenal hyperplasia (CAH) due to 21-hydroxylase deficiency aged 0-3 years: Update on a retrospective multicentre analysis using the I-CAH registry**
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- LI. **Follow-up of individuals with gender identity disorders: A long and challenging process**
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- LII. **Contemporary surgical approach in CAH 46XX-Results from the I-DSD/I-CAH Registries**
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- LIII. **Development Of An International Benchmark For Sick Day Episodes As A Core Clinical Outcome In People With Congenital Adrenal Hyperplasia**
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- LIV. **Global Practice of Glucocorticoid and Mineralocorticoid Treatment in Children and Adults with Congenital Adrenal Hyperplasia - Insights from the I-CAH Registry**
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- LV. **Two siblings with hypophosphatemic rickets: SLC34A3 gene mutations with different clinical phenotypes**
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- LVI. **Long-Term Outcome In Leydig Cell Hypoplasia**
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- LVII. **Characteristics of puberty, pubertal height gain and final height in children with classical 21 hydroxylase deficiency**
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- LVIII. **The investigation of genetic etiology in familial cases with congenital hypothyroidism**
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- LIX. **Targeted Panel Gene Sequencing for Identification of Genetic Etiology of 46, XY Disorders of Sex Development**
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Metrics

Publication: 377

Citation (WoS): 3191

Citation (Scopus): 3461

H-Index (WoS): 27

H-Index (Scopus): 30