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Kişisel Bilgiler

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SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

I. **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., Helvacioglu D., HALİLOĞLU B., KAYGUSUZ S. B., Yavas Abali Z., SEVEN MENEVŞE T., et al.
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II. **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, cilt.36, sa.4, ss.401-408, 2023 (SCI-Expanded)

III. **Etiological analysis of hypophosphatemia: A single-center experience**
Eltan M., Alavanda C., Abali Z. Y., Bayramoglu E., Kaygusuz S. B., Helvacioglu D., Tosun B. G., Menevse T. S., Ata P., Guran T., et al.
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IV. **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.
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V. **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, cilt.16, sa.SUPPL 1, ss.61-62, 2022 (SCI-Expanded)

VI. **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., Gökce İ., Abali S., Alavanda C., Arman A., Kırkgöz T., Güran T., Hatun S., et al.
CALCIFIED TISSUE INTERNATIONAL, cilt.110, sa.4, ss.441-450, 2022 (SCI-Expanded)

VII. **Biallelic PPP2R3C mutations are associated with partial and complete gonadal dysgenesis in 46,XY and 46,XX individuals**
Çiçek D., Warr N., Yesil G., Eker H. K., Bas F., Poyrazoglu S., Darendeliler F., Direk G., Hatipoğlu N., Eltan M., et al.
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VIII. **A rare cause of hypercalcemia: Congenital Lactase Deficiency**
Eltan M., Alavanda C., Abalı S., Abali Z. Y., Kaygusuz S. B., Gürpınar Tosun B., Seven Menevşe T., Helvacioglu D., Güran T., Ata P., et al.
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IX. **46,XY DSD due to biallelic DHX37 gene mutations**
Eltan M., Helvacioglu D., Ates E. A., Abali Z. Y., Demircioğlu S., Bereket A., Güran T.
HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.393-394, 2021 (SCI-Expanded)

X. **Correlation of 11-oxygenated C19 androgens with the clinical and biochemical characteristics in**

premature adrenarche

Abali Z. Y., Eltan M., Helvacioglu D., Yaman A., Demircioğlu S., Bereket A., Güran T.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.120-121, 2021 (SCI-Expanded)

XI. **Acetazolamide treatment in a patient with pseudohypoparathyroidism with venous calcification**

Seven Menevşe T., Gürpınar Tosun B., Abali Z. Y., Helvacioglu D., Kaygusuz S. B., Eltan M., Buğdaycı O., Güran T., Bereket A., Demircioğlu S.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.75, 2021 (SCI-Expanded)

XII. **Does Genotype–Phenotype Correlation Exist in Vitamin D-Dependent Rickets Type IA: Report of 13 New Cases and Review of the Literature**

Kaygusuz S. B., Alavanda C., Kırkgöz T., Eltan M., Yavas Abali Z., Helvacioglu D., Güran T., Ata P., Bereket A., Demircioğlu S.

Calcified Tissue International, cilt.108, sa.5, ss.576-586, 2021 (SCI-Expanded)

XIII. **Cinacalcet as a First-Line Treatment in Neonatal Severe Hyperparathyroidism Secondary to Calcium Sensing Receptor (CaSR) Mutation**

Gulcan-Kersin S., Kirkgoz T., Eltan M., Rzayev T., Ata P., Bilgen H. S., Özek E., Bereket A., Turan S.

HORMONE RESEARCH IN PAEDIATRICS, cilt.93, sa.5, ss.313-321, 2020 (SCI-Expanded)

XIV. **A Rare Cause of Hypophosphatemia: Raine Syndrome Changing Clinical Features with Age**

Eltan M., Alavanda C., Yavas Abali Z., Ergenekon P., Yalindag Ozturk N., Sakar M., Dağçınar A., Kırkgöz T., Kaygusuz S. B., Gökdemir Y., et al.

CALCIFIED TISSUE INTERNATIONAL, cilt.107, ss.96-103, 2020 (SCI-Expanded)

XV. **Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation**

Eltan M., Yavas Abali Z., Arslan Ates E., Kırkgöz T., Kaygusuz S. B., Türkyllmaz A., Bereket A., Turan S., Güran T.

Hormone Research in Paediatrics, cilt.92, sa.4, ss.262-268, 2020 (SCI-Expanded)

XVI. **Cushing Syndrome due to an adrenacortical carcinoma in a baby with atypical Beckwith-Wiedemann Syndrome**

Eltan M., Cerit K., Kaygusuz S. B., Ates E., Eker N., Bagci P., Ergelen R., Turan S., Bereket A., Güran T.

HORMONE RESEARCH IN PAEDIATRICS, cilt.91, ss.371, 2019 (SCI-Expanded)

XVII. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**

Kaygusuz S. B., Arman A., Abalı S., Ata P., Kırkgöz T., Eltan M., Abali Z. Y., Helvacioglu D., Tosun B. G., Menevse T. S., et al.

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XVIII. **A Real-Life Experience with A New Insulin Co-Formulation Degludec/Aspart For One Year In Poorly Controlled Children And Adolescents With Type 1 Diabetes**

Kırkgöz T., Eltan M., Kaygusuz S. B., Abali Z. Y., Güran T., Bereket A., Turan S.

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XIX. **A Case Of Syndromic Hypopituitarism**

Kaygusuz S. B., Ates E. A., Kırkgöz T., Eltan M., Abali Z. Y., Helvacioglu D., Menevse T. S., Tosun B. G., Tutar E., Volkan B., et al.

HORMONE RESEARCH IN PAEDIATRICS, cilt.91, ss.589, 2019 (SCI-Expanded)

XX. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**

Kaygusuz S. B., Ata P., Kırkgöz T., Abali Z. Y., Eltan M., Tosun B. G., Menevse T. S., Helvacioglu D., Güran T., Arman A., et al.

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XXI. **Evaluation of brain MRI lesions in 381 girls with central precocious puberty**

Helvacioglu D., GÜRAN T., KIRKGÖZ T., Atay Z., Abali Z. Y., ELTAN M., KAYGUSUZ S. B., Seven T., Gurpinar B., Turan S., et al.

HORMONE RESEARCH IN PAEDIATRICS, cilt.91, ss.329-330, 2019 (SCI-Expanded)

XXII. **Simplifying the interpretation of steroid metabolome data by a machine-learning approach**

Kırkgöz T., Kilic S., Abali Z. Y., Yaman A., Kaygusuz S. B., Eltan M., Turan S., Haklar G., Sagiroglu M. S., Bereket A., et al.

HORMONE RESEARCH IN PAEDIATRICS, cilt.91, ss.128, 2019 (SCI-Expanded)

XXIII. **A rare cause of hypophosphatemia: Raine Syndrome**
Eltan M., Ata P., Kırkgöz T., Alavanda C., Kaygusuz S. B., Menevse T. S., Tosun B. G., Abali Z. Y., Helvacioglu D., Güran T., et al.
HORMONE RESEARCH IN PAEDIATRICS, cilt.91, ss.384, 2019 (SCI-Expanded)

XXIV. **Evaluation of growth and puberty in a child with a novel TBX19 gene mutation and review of the literature**
Abali Z. Y., Yeşil G., Kırkgöz T., Kaygusuz S. B., Eltan M., Turan S., Bereket A., Güran T.
HORMONES-INTERNATIONAL JOURNAL OF ENDOCRINOLOGY AND METABOLISM, cilt.18, sa.2, ss.229-236, 2019 (SCI-Expanded)

Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

I. **Genotype and Phenotype Characterization ofTurkish Patients with Vitamin D Dependent RicketsType IA**
KAYGUSUZ S. B., ATA P., KIRKGÖZ T., YAVAŞ ABALI Z., ELTAN M., GÜRPINAR T. B., SEVEN M. T., HELVACIOĞLU D., ARMAN A., GÜRAN T., et al.
European Society for Paediatric Endocrinology (ESPE)58th Annual Meeting, Vienna, VİYANA, Avusturya, 19 - 21 Eylül 2019, cilt.91

II. **A rare cause of hypophosphatemia: Raine Syndrome**
ELTAN M., ATA P., KIRKGÖZ T., ALAVANDA C., KAYGUSUZ S. B., SEVEN M. T., GURPINAR TOSUN B., YAVAŞ ABALI Z., GÜRAN T., ELÇİOĞLU H. N., et al.
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III. **Yenidoğanda Nadir Bir Hiperkalsemi nedeni : Kalsiyum Duyarlı Reseptör Mutasyonuna bağlı Ağır neonatal Hiperparatiroidizm**
Rzayev T., KIRKGÖZ T., Özdemir H., ATA P., BİLGEN H. S., MEMİŞOĞLU A., ELTAN M., ÖZEK E., Turan S.
27. Ulusal Neonatoloji Kongresi, Antalya, Türkiye, 3 - 07 Nisan 2019

IV. **Segmental Aşırı büyüme kliniği olan olguda somatik PIK3CA mutasyonu**
Yavaş Abalı Z., Arslan Ateş E., Türkyılmaz A., Salman A., Kırkgöz T., Kaygusuz S. B., Eltan M., Turan S., Bereket A., Güran T.
3. Ege Endokrin Hastalıklar ve Genetik Sempozyumu, İzmir, Türkiye, 7 - 09 Mart 2019

V. **A further case of autosomal recessive brachyolmia having a novel mutation in PAPSS2 gene**
Arslan Ateş E., Eltan M., Türkyılmaz A., Alavanda C., Söylemez M. A., Geçkinli B. B., Güney A. İ., Güran T., Arman A.
13 ULUSAL TIBBİ GENETİK KONGRESİ, Antalya, Türkiye, 7 - 11 Kasım 2018

Metrikler

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H-İndeks (Scopus): 4