

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., Helvacioğlu D., HALİLOĞLU B., KAYGUSUZ S. B., Yavas Abali Z., SEVEN MENEVŞE T., et al.
HORMONE RESEARCH IN PAEDIATRICS, cilt.96, sa.5, ss.527-537, 2023 (SCI-Expanded)
- II. **Molecular analysis of MKRN3 gene in Turkish girls with sporadic and familial idiopathic central**
KIRKGÖZ T., KAYGUSUZ S. B., ALAVANDA C., Helvacioğlu 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. **Rare forms of congenital adrenal hyperplasia**
GÜRPINAR TOSUN B., GÜRAN T.
Clinical Endocrinology, 2023 (SCI-Expanded)
- IV. **Glucagon response to hypoglycemia during extended oral glucose tolerance test in children with cystic fibrosis and comparing with healthy peers**
HALİLOĞLU B., SEVEN MENEVŞE T., GÜRPINAR TOSUN B., GÜRAN T., DEMİRCİOĞLU S., Ispir T., GÖKDEMİR Y., ERDEM ERALP E., BEREKET A.
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.199-200, 2022 (SCI-Expanded)
- V. **Wide phenotypical spectrum with the same karyotype: Mixed gonadal dysgenesis**
Seven Menevşe T., Gürpınar Tosun B., Helvacioğlu D., Abali Z. Y., Kirmizibekmez H., Dursun F., Demircioğlu S., Bereket A., Güran T.
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.566, 2022 (SCI-Expanded)
- VI. **Diagnostic Features and Risk Factors for Childhood Thyroid Cancers**
ŞAHİN P., GÜRPINAR TOSUN B., YUMUŞAKHUYLU A. C., GÜRAN T., Helvacioğlu D., Abali Z. Y., HALİLOĞLU B., OYSU Ç., BEREKET A., DEMİRCİOĞLU S.
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.407, 2022 (SCI-Expanded)
- VII. **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, cilt.95, sa.SUPPL 2, ss.377-378, 2022 (SCI-Expanded)
- VIII. **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., Helvacioğlu D., Gürpınar Tosun B., Haliloğlu B., et al.
HORMONE RESEARCH IN PAEDIATRICS, cilt.95, sa.SUPPL 2, ss.364, 2022 (SCI-Expanded)
- IX. **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, cilt.95, sa.SUPPL 2, ss.139, 2022 (SCI-Expanded)
- X. **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, cilt.95, sa.SUPPL 2, ss.117-118, 2022 (SCI-Expanded)

XI. **Gonadal Outcome in 17beta-HSD deficiency and 5alpha-reductase deficiency**

Boogers L. S., Bruggenwirth H. T., van Bever Y., Hersmus R., Bryce J., Ahmed S. F., Lucas-Herald A. K., Baronio F., Cools M., Ellaithi M., et al.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.91-92, 2021 (SCI-Expanded)

XII. **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.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.92, 2021 (SCI-Expanded)

XIII. **Clinical and hormonal evolution of aldosterone synthase deficiency: Is complete remission possible?**

Gürpınar Tosun B., Demirkol Y. K., Seven Menevşe T., Kaygusuz S. B., Ozbek M. N., Altıncık S. A., Mammadova J., Cayir A., Döğ er E., Bayramoglu E., et al.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.184-185, 2021 (SCI-Expanded)

XIV. **Adrenocortical hormone profiles do not predict the molecular etiology in non-CAH primary adrenal insufficiency**

Seven Menevşe T., Demirkol Y. K., Gürpınar Tosun B., Bayramoglu E., Yildiz M., Acar S., Karaca S. E., Orbak Z., Onder A., Sobu E., et al.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.64-65, 2021 (SCI-Expanded)

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

Seven Menevşe T., Gürpınar Tosun B., Abalı Z. Y., Helvacioğlu D., Kaygusuz S. B., Eltan M., Buğ daycı O., Güran T., Bereket A., Demircioğlu S.

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

XVI. **A rare cause of hypercalcemia: Congenital Lactase Deficiency**

Eltan M., Alavanda C., Abalı S., Abalı Z. Y., Kaygusuz S. B., Gürpınar Tosun B., Seven Menevşe T., Helvacioğlu D., Güran T., Ata P., et al.

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

XVII. **Hyperinsulinemic hypoglycemia due to biallelic mutations in the DNAJC3 gene**

Gürpınar Tosun B., Seven Menevşe T., Esen N., Demircioğlu S., Yesilyurt A., Güran T., Bereket A.

HORMONE RESEARCH IN PAEDIATRICS, cilt.94, sa.SUPPL 1, ss.107, 2021 (SCI-Expanded)

Diğer Dergilerde Yayınlanan Makaleler

I. **Dysosteosclerosis: Clinical and Radiological Evolution Reflecting Genetic Heterogeneity**

DEMİRCİOĞ LU 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, cilt.6, sa.8, 2022 (ESCI)

Metrikler

Yayın: 19

Atıf (WoS): 2

Atıf (Scopus): 2

H-İndeks (WoS): 1

H-İndeks (Scopus): 1