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Published journal articles indexed by SCI, SSCI, and AHCI

- 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, vol.96, no.5, pp.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, vol.36, no.4, pp.401-408, 2023 (SCI-Expanded)
- III. **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, vol.95, no.SUPPL 2, pp.199-200, 2022 (SCI-Expanded)
- IV. **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, vol.95, no.SUPPL 2, pp.566, 2022 (SCI-Expanded)
- 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, vol.16, no.SUPPL 1, pp.61-62, 2022 (SCI-Expanded)
- VI. **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., Döğler E., Bayramoğlu E., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.184-185, 2021 (SCI-Expanded)
- VII. **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., Helvacioğlu D., Güran T., Ata P., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.75, 2021 (SCI-Expanded)
- VIII. **Acetazolamide treatment in a patient with pseudohypoparathyroidism with venous calcification**
Seven Menevşe T., Gürpınar Tosun B., Abali Z. Y., Helvacioğlu D., Kaygusuz S. B., Eltan M., Buğdaycı O., Güran T., Bereket A., Demircioğlu S.
HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.75, 2021 (SCI-Expanded)
- IX. **Adrenocortical hormone profiles do not predict the molecular etiology in non-CAH primary adrenal insufficiency**
Seven Menevşe T., Demirkol Y. K., Gürpınar Tosun B., Bayramoğlu E., Yıldız M., Acar S., Karaca S. E., Orbak Z., Onder

A., Sobu E., et al.

HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.64-65, 2021 (SCI-Expanded)

- X. **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, vol.94, no.SUPPL 1, pp.92, 2021 (SCI-Expanded)

- XI. **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, vol.94, no.SUPPL 1, pp.107, 2021 (SCI-Expanded)

- XII. **Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study**

Groeneweg S., Van Geest F. S., Abacı A., Alcantud A., Ambegaonkar G. P., Armour C. M., Bakhtiani P., Barca D., Bertini E. S., Van Beynum I. M., et al.

LANCET DIABETES & ENDOCRINOLOGY, vol.8, no.7, pp.594-605, 2020 (SCI-Expanded)

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