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Articles Published in Journals That Entered SCI, SSCI and AHCI Indexes

I. **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., DAĞÇINAR A., KIRKGÖZ T., KAYGUSUZ S. B. , GÖKDEMİR Y., et al.
CALCIFIED TISSUE INTERNATIONAL, vol.107, pp.96-103, 2020 (Journal Indexed in SCI)

II. **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., KIRKGÖZ T., KAYGUSUZ S. B. , Türkyllmaz A., BEREKET A., Turan S., GÜRAN T.
Hormone Research in Paediatrics, vol.92, pp.262-268, 2020 (Journal Indexed in SCI)

III. **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., ÖZEK E., BEREKET A., Turan S.
Hormone Research in Paediatrics, 2020 (Journal Indexed in SCI Expanded)

IV. **A Case Of Syndromic Hypopituitarism**
KAYGUSUZ S. B. , Ates E. A. , KIRKGÖZ T., ELTAN M., Abali Z. Y. , Helvacioglu D., Menevse T. S. , Tosun B. G. , Tutar E., Volkan B., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.589, 2019 (Journal Indexed in SCI)

V. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**
KAYGUSUZ S. B. , ATA P., KIRKGÖZ T., Abali Z. Y. , ELTAN M., Tosun B. G. , Menevse T. S. , Helvacioglu D., GÜRAN T., ARMAN A., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.215, 2019 (Journal Indexed in SCI)

VI. **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, vol.91, pp.329-330, 2019 (Journal Indexed in SCI)

VII. **Simplifying the interpretation of steroid metabolome data by a machine-learning approach**
KIRKGÖZ 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, vol.91, pp.128, 2019 (Journal Indexed in SCI)

VIII. **A rare cause of hypophosphatemia: Raine Syndrome**
ELTAN M., ATA P., KIRKGÖZ 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, vol.91, pp.384, 2019 (Journal Indexed in SCI)

IX. **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, vol.91, pp.371, 2019 (Journal Indexed in SCI)

X. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**
KAYGUSUZ S. B. , ARMAN A., ABALI S., ATA P., KIRKGÖZ T., ELTAN M., Abali Z. Y. , Helvacioglu D., Tosun B. G. ,

Menevse T. S. , et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.214, 2019 (Journal Indexed in SCI)

XI. **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**
KIRKGÖZ T., ELTAN M., KAYGUSUZ S. B. , Abali Z. Y. , GÜRAN T., BEREKET A., Turan S.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.534-535, 2019 (Journal Indexed in SCI)

XII. **Evaluation of growth and puberty in a child with a novel TBX19 gene mutation and review of the literature**
Abali Z. Y. , YEŞİL G., KIRKGÖZ T., KAYGUSUZ S. B. , ELTAN M., Turan S., BEREKET A., GÜRAN T.
HORMONES-INTERNATIONAL JOURNAL OF ENDOCRINOLOGY AND METABOLISM, vol.18, pp.229-236, 2019 (Journal Indexed in SCI)

Refereed Congress / Symposium Publications in Proceedings

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, Austria, 19 - 21 September 2019, vol.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.
European Society for Paediatric Endocrinology (ESPE)58th Annual Meeting, Vienna, VİYANA, Austria, 19 - 21 September 2019, vol.91

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, Turkey, 3 - 07 April 2019

IV. **Segmental Aşırı büyüme kliniği olan olguda somatik PIK3CA mutasyonu**
YAVAŞ ABALI Z., Arslan Ateş E., türkyılmaz a., SALMAN A., KIRKGÖZ T., KAYGUSUZ S. B. , ELTAN M., Turan S., BEREKET A., GÜRAN T.
3. Ege Endokrin Hastalıklar ve Genetik Sempozyumu, Turkey, 7 - 09 March 2019

V. **A further case of autosomal recessive brachyolmia having a novel mutation in PAPSS2 gene**
ARSLAN ATEŞ E., ELTAN M., TÜRKYILMAZ A., ALAVANDA C., SÖYLEMEZ M. A. , GEÇKİNLİ B. B. , GÜNEY A. İ. , GÜRAN T., ARMAN A.
13 ULUSAL TIBBİ GENETİK KONGRESİ, Turkey, 7 - 11 November 2018

Citations

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