

Res. Asst. SARE BETÜL KAYGUSUZ

Personal Information

Email: skaygusuz@marmara.edu.tr

Web: <https://avesis.marmara.edu.tr/skaygusuz>

Education Information

Expertise In Medicine, Bezm-İ Âlem Vakıf Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Turkey 2012 - Continues
Undergraduate, İstanbul Üniversitesi, İstanbul Tıp Fakültesi, Turkey 2006 - 2012

Academic Titles / Tasks

Research Assistant, Bezm-İ Âlem Vakıf Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2012 - Continues

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

- I. **Is quail egg a potential endocrine disrupter?**
SÜREKLİ KARAKUŞ Ö., ARABACI TAMER S., Levent H. N. , KAYGUSUZ S. B. , DEMİRCİOĞLU S., AKAKIN D., GÜRAN T., YEGEN B., BEREKET A.
NEUROENDOCRINOLOGY, vol.111, pp.32, 2021 (Journal Indexed in SCI)
- II. **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., Helvacioğlu D., Güran T., Ata P., Bereket A., Demircioğlu S.
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- III. **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., Dağcınar A., Kırkgöz T., Kaygusuz S. B. , Gökdemir Y., et al.
CALCIFIED TISSUE INTERNATIONAL, vol.107, pp.96-103, 2020 (Journal Indexed in SCI)
- IV. **Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation**
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- V. **A Case Of Syndromic Hypopituitarism**
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- VI. **A rare cause of hypophosphatemia: Raine Syndrome**
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- VII. **Simplifying the interpretation of steroid metabolome data by a machine-learning approach**
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- VIII. **Evaluation of brain MRI lesions in 381 girls with central precocious puberty**
Helvacioğlu D., GÜRAN T., KIRKGÖZ T., Atay Z., Abali Z. Y., ELTAN M., KAYGUSUZ S. B., Seven T., Gürpınar B., Turan S., et al.
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- IX. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**
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- X. **Cushing Syndrome due to an adrenocortical carcinoma in a baby with atypical Beckwith-Wiedemann Syndrome**
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HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.371, 2019 (Journal Indexed in SCI)
- XI. **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., Helvacioğlu D., Tosun B. G., Menevse T. S., et al.
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- XII. **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)
- XIII. **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, no.2, pp.229-236, 2019 (Journal Indexed in SCI)
- XIV. **An Unusual Cause of Short Stature**
KAYGUSUZ S. B., Atay Z., KIRKGÖZ T., GÜRAN T., BEREKET A., Turan S.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.184-185, 2018 (Journal Indexed in SCI)
- XV. **Recurrent Hypoglycemia in a Preschooler Girl with Overgrowth: Isolated ACTH-Deficiency with a Novel TPIT Mutation**
Abali Z. Y., Yesil G., KIRKGÖZ T., KAYGUSUZ S. B., Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.132, 2018 (Journal Indexed in SCI)
- XVI. **Isolated Congenital Central Hypothyroidism due to a Novel Mutation in TSH Beta Subunit Gene**
KIRKGÖZ T., ÖZHAN B., Cetin O., KAYGUSUZ S. B., Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.597, 2018 (Journal Indexed in SCI)
- XVII. **An 18 Month Old Boy with Hypoglycemic Convulsion and Obesity Due to POMC Deficiency**
KAYGUSUZ S. B., Yesil G., KIRKGÖZ T., Turan S., BEREKET A., GÜRAN T.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.514, 2018 (Journal Indexed in SCI)

Refereed Congress / Symposium Publications in Proceedings

- I. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**
KAYGUSUZ S. B., ARMAN A., ABALI S., ATA P., KIRKGÖZ T., YAVAŞ ABALI Z., HELVACIOĞLU D., GÜRPINAR T. B., SEVEN M. T., GÜRAN T., et al.
European Society for Paediatric Endocrinology (ESPE) 58th Annual Meeting, Vienna, VIYANA, Austria, 19 - 21 September 2019
- 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. Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type 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.

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IV. Segmental Aşırı büyüme kliniği olan olguda somatik PIK3CA mutasyonu

YAVAŞ ABALI Z., Arslan Ateş E., türkylmaz 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. An 18 Month Old Boy with Hypoglycemic Convulsion and Obesity Due to POMC Deficiency

KAYGUSUZ S. B. , YEŞİL G., KIRKGÖZ T., DEMİRCİOĞLU S., BEREKET A.

HORMONE RESEARCH IN PAEDIATRICS-57th Annual Meeting of the European Society for Paediatric Endocrinology, 27 - 29 September 2018, vol.90, pp.514

VI. Isolated Congenital Central Hypothyroidism due to a Novel Mutation in TSH Beta Subunit Gene

KIRKGÖZ T., ÖZHAN B., ÇETİN G. O. , KAYGUSUZ S. B. , DEMİRCİOĞLU S., BEREKET A., GÜRAN T.

HORMONE RESEARCH IN PAEDIATRICS-57th Annual Meeting of the European Society for Paediatric Endocrinology, 27 - 29 September 2018, vol.90, pp.597

VII. Recurrent Hypoglycemia in a Preschooler Girl with Overgrowth: Isolated ACTH-Deficiency with a Novel TPIT Mutation

YAVAŞ ABALI Z., YEŞİL G., KIRKGOZ T., KAYGUSUZ S. B. , DEMİRCİOĞLU S., BEREKET A., GÜRAN T.

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Citations

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