# Res. Asst. PhD İHSAN TURAN

## Personal Information

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## Education Information

Expertise In Medicine, Marmara University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Turkey 2010 - 2014
Under Graduate, Istanbul University, Cerrahpaşa Tıp Fakültesi, Turkey 2002 - 2008

## Foreign Languages

English, C1 Advanced

## Dissertations

Expertise In Medicine, Bronşiolitis Obliteranslı Hastalarda Uyku Bozuklukları, Marmara University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkarı, 2014

## Research Areas

Medicine, Health Sciences, Internal Medicine Sciences, Child Health and Diseases, Pediatric Endocrinology and Metabolism

## Academic Titles / Tasks

Research Assistant PhD, Cukurova University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, 2015 - Continues

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

I. **21-Hydroxylase deficiency: Mutational spectrum and Genotype-Phenotype relations analyses by next-generation sequencing and multiplex ligation-dependent probe amplification**
Turan I., Tastan M., Boga D. D. , GÜRBÜZ F., KOTAN L. D. , TULİ A., YÜKSEL B.
EUROPEAN JOURNAL OF MEDICAL GENETICS, vol.63, 2020 (Journal Indexed in SCI)

II. **An infant with hyponatremia, hyperkalemia, and metabolic acidosis associated with urinary tract infection: Answers**
ATMIŞ B., TURAN İ., MELEK E., KARABAY BAYAZIT A.
PEDIATRIC NEPHROLOGY, vol.34, pp.1739-1741, 2019 (Journal Indexed in SCI)

III. **An infant with hyponatremia, hyperkalemia, and metabolic acidosis associated with urinary tract infection: Questions**
ATMIŞ B., TURAN İ., MELEK E., KARABAY BAYAZIT A.
PEDIATRIC NEPHROLOGY, vol.34, pp.1737, 2019 (Journal Indexed in SCI)

IV. **BMP4 mutations as a novel cause of normosmic hypogonadotropic hypogonadism**

Topaloglu A. K. , Yildirim R., KOTAN L. D. , Akkus G., Unal E., Turan I., Dilek S., Tastan M., GÜRBÜZ F., YÜKSEL B.
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.113-114, 2019 (Journal Indexed in SCI)

V. **Determination of cerebral edema with serial measurement of optic nerve sheath diameter during treatment in children with diabetic ketoacidosis: a longitudinal study**
KENDIR O. T. , YILMAZ H. L. , ÖZKAYA A. K. , TURAN İ., GOKAY S. S. , BILEN S., YILDIZDAŞ R. D. , YÜKSEL B.
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.32, pp.943-949, 2019 (Journal Indexed in SCI)

VI. **Unusual and early onset IPEX syndrome: a case report**
DOĞRUEL D., GÜRBÜZ F., TURAN İ., ALTINTAŞ D. U. , YILMAZ M., YÜKSEL B.
TURKISH JOURNAL OF PEDIATRICS, vol.61, pp.580-584, 2019 (Journal Indexed in SCI)

VII. **Efficiency of Single Dose of Tolvaptan Treatment During the Triphasic Episode After Surgery for Craniopharyngioma**
GÜRBÜZ F., Tastan M., TURAN İ., YÜKSEL B.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.11, pp.202-206, 2019 (Journal Indexed in SCI)

VIII. **Prevalence and associated phenotypes of PLXNA1 variants in normosmic and anosmic idiopathic hypogonadotropic hypogonadism**
KOTAN L. D. , Isik E., TURAN İ., Mengen E., AKKUŞ G., Tastan M., GÜRBÜZ F., YÜKSEL B., TOPALOĞLU A. K.
CLINICAL GENETICS, vol.95, pp.320-324, 2019 (Journal Indexed in SCI)

IX. **ATTENTION TO TRANSIENT PSEUDOHYPOALDOSTERONISM IN INFANCY WITH PYELONEPHRITIS**
Sapmaz M., Sahin G., TURAN İ., ATMIŞ B., MELEK E., KARABAY BAYAZIT A.
PEDIATRIC NEPHROLOGY, vol.33, pp.1967, 2018 (Journal Indexed in SCI)

X. **The Authors' Reply: In systemic pseudohypoaldosteronism type 1 skin manifestations are not rare and the disease is not transient**
TURAN İ., TOPALOĞLU A. K. , YÜKSEL B.
CLINICAL ENDOCRINOLOGY, vol.89, pp.241-242, 2018 (Journal Indexed in SCI)

XI. **Molecular genetic studies in a case series of isolated hypoaldosteronism due to biosynthesis defects or aldosterone resistance.**
TURAN İ., Kotan L. D. , TASTAN M., GURBUZ F., TOPALOĞLU A. K. , YÜKSEL B.
Clinical endocrinology, vol.88, pp.799-805, 2018 (Journal Indexed in SCI)

XII. **A Turkish Family with 46,XY Disorder of Sex Development Due to 17b-Hydroxysteroid Dehydrogenase Type 3 Deficiency**
GÜRBÜZ F., TURAN İ., Tastan M., YÜKSEL B.
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.582, 2018 (Journal Indexed in SCI)

XIII. **Hypogonadotropic Hypogonadism due to Novel FGFR1 Mutations**
AKKUŞ G., KOTAN L. D. , Durmaz E., Mengen E., TURAN İ., Ulubay A., GÜRBÜZ F., YÜKSEL B., Tetiker T., TOPALOĞLU A. K.
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.9, pp.95-100, 2017 (Journal Indexed in SCI)

XIV. **CCDC141 Mutations in Idiopathic Hypogonadotropic Hypogonadism**
TURAN İ., Hutchins B. I. , Hacihamdioglu B., KOTAN L. D. , GÜRBÜZ F., Ulubay A., Mengen E., YÜKSEL B., Wray S., TOPALOĞLU A. K.
JOURNAL OF CLINICAL ENDOCRINOLOGY & METABOLISM, vol.102, pp.1816-1825, 2017 (Journal Indexed in SCI)

XV. **A CASE OF CONGENITAL GENERALIZED LIPODYSTROPHY TYPE 2 WITH NOVEL BSCL2 GENE MUTATION**
GÜRBÜZ F., TURAN İ., Tastan M., TOPALOĞLU A. K. , YÜKSEL B.
HORMONE RESEARCH IN PAEDIATRICS, vol.88, pp.141-142, 2017 (Journal Indexed in SCI)

XVI. **A RARE CAUSE OF CONGENITAL ADRENAL HYPERPLASIA: CONGENITAL LIPOID ADRENAL HYPERPLASIA**
GÜRBÜZ F., TURAN İ., TOPALOĞLU A. K. , YÜKSEL B.
HORMONE RESEARCH IN PAEDIATRICS, vol.88, pp.566, 2017 (Journal Indexed in SCI)

XVII. **Idiopathic Hypogonadotropic Hypogonadism Caused by Inactivating Mutations in SRA1.**

Kotan L. D. , COOPER C., DARCAN Ş., CARR I., ÖZEN S., YAN Y., HAMEDANI M., Gürbüz F., MENGEN E., TURAN İ., et al.
Journal of clinical research in pediatric endocrinology, vol.8, pp.125-34, 2016 (Journal Indexed in SCI)

XVIII. **Prevalence of Neisseria meningitidis carriage: a small-scale survey in Istanbul, Turkey**
Kadayifci E. K. , Merdan D. G. , Soysal A., Karaaslan A., Atici S., Durmaz R., BORAN P., TURAN İ., Soyletir G., Bakir M.
JOURNAL OF INFECTION IN DEVELOPING COUNTRIES, vol.10, pp.413-417, 2016 (Journal Indexed in SCI)

XIX. **Sleep disordered breathing and sleep quality in children with bronchiolitis obliterans**
Uyan Z. S. , TURAN İ., Ay P., ÇAKIR E., Ozturk E., Gedik A. H. , GÖKDEMİR Y., Erdem E., Sen V., KARADAĞ B. T. , et al.
PEDIATRIC PULMONOLOGY, vol.51, pp.308-315, 2016 (Journal Indexed in SCI)

XX. **Idiopathic Hypogonadotrophic Hypogonadism Caused by Inactivating Mutations in SRA1**
KOTAN L. D. , Cooper C., DARCAN Ş., Carr I., ÖZEN S., Yan Y., Hamedani M. K. , GÜRBÜZ F., Mengen E., TURAN İ., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.86, pp.58-59, 2016 (Journal Indexed in SCI)

XXI. **Inactivating Mutations in CCDC141 Causing Idiopathic Hypogonadotrophic Hypogonadism/Kallmann Syndrome**
TURAN İ., Hutchins B. I. , Hacihamdioglu B., Ozbek M. N. , KOTAN L. D. , ÖZKAN Y., Stoner H., Cheng P. J. , GÜRBÜZ F., Mengen E., et al.
HORMONE RESEARCH IN PAEDIATRICS, vol.86, pp.58, 2016 (Journal Indexed in SCI)

XXII. **Complete idiopathic hypogonadotropic hypogonadism due to homozygous GNRH1 mutations in the mutational hot spots in the region encoding the decapeptide**
MENGEN E., TUNC S., KOTAN L. D. , NALBANTOGLU O., Demir K., GÜRBÜZ F., TURAN İ., ŞEKER G., YÜKSEL B., TOPALOĞLU A. K.
Hormone Research in Paediatrics, vol.85, pp.107-111, 2016 (Journal Indexed in SCI)

## Articles Published in Other Journals

I. **Urolithiasis Frequency and Risk Factors in Home Ventilated Patients with Tracheostomy**
tolunay i., yıldıztaş d., horoz ö. ö. , MELEK E., ATMIŞ B., YÜKSEL B., TURAN İ.
Turkish Journal of Pediatric Emergency and Intensive Care Medicine, pp.92-95, 2017 (Refereed Journals of Other Institutions)

II. **Ev Tipi Ventilatör ile İzlenen Trakeostomili Hastalarda Ürolitiazis Sıklığı ve Risk Faktörleri**
Tolunay İ., Yıldızdaş R. D. , Horoz Ö. Ö. , MELEK E., Atmış B., Yüksel B., Turan İ.
J Pediatr Emerg Intensive Care Med, vol.4, pp.92-95, 2017 (National Refreed University Journal)

## Refereed Congress / Symposium Publications in Proceedings

I. **Konjenital Adrenal Hiperplazi Tanılı Hastalarda Hipertansiyon**
TURAN İ.
4. Güney İlleri Çocuk Endokrinolojisi Toplantısı, Adana, Turkey, 02 December 2017

II. **LHCGR Gen Mutasyonuna Bağlı46, XY Yetersiz Virilize Olgu Sunumu**
TURAN İ.
4.Güney İlleri Çocuk Endokrinolojisi Toplantısı, Turkey, 02 December 2017

III. **Hipokalsemik Hastada CYP27B1 Mutasyonuna Bağlı Dirençli Rikets Olgusu**
TAŞTAN M., GÜRBÜZ F., TURAN İ., YÜKSEL B.
4. Güney İlleri Çocuk Endokrinolojisi Toplantısı, Turkey, 02 December 2017

IV. **17-Beta-OHSD Eksikliğine Bağlı Ambiygus Genitale**
GÜRBÜZ F., TURAN İ., TAŞTAN M., YÜKSEL B.
4. Güney İlleri Çocuk Endokrinolojisi Toplantısı, Turkey, 02 December 2017

V. **Novel LHCGR Gen Mutasyonuna Bağlı 46,XY Yetersiz Virilizasyon**
TURAN İ., GÜRBÜZ F., TAŞTAN M., KOTAN GEDİK L. D. , YÜKSEL B.
4. Güney İlleri Çocuk Endokrinolojisi Toplantısı, Turkey, 02 December 2017

VI. **A CASE OF CONGENITAL GENERALIZED LIPODYSTROPHY TYPE 2 WITH NOVEL BSCL2 GENE MUTATION**
GÜRBÜZ F., TURAN İ., TAŞTAN M., TOPALOĞLU A. K. , YÜKSEL B.
10th Joint Meeting of Paediatric Endocrinology, PES-APEG-APPES-ASPAE-CSPEM-ESPE-JSPE-SLEP, Washington., Turkey, 14 - 19 September 2017, vol.88, pp.1-628

VII. **A RARE CAUSE OF CONGENITAL ADRENAL HYPERPLASIA:CONGENITAL LIPOID ADRENAL HYPERPLASIA**
GÜRBÜZ F., TURAN İ., TOPALOĞLU A. K. , YÜKSEL B.
10. International Meeting of Pediatric Endocrinology, Washington, Kiribati, 14 - 17 September 2017

VIII. **İdiyopatik Hipogonadotropik Hipogonadizm'de CCDC141 Mutasyonları**
TURAN İ., HACIHAMDİOĞLU B., KOTAN GEDİK L. D. , TAŞTAN M., GÜRBÜZ F., YÜKSEL B., TOPALOĞLU A. K.
XXI. ULUSAL PEDİATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Turkey, 26 - 30 April 2017

IX. **Nadir bir Kongenital Adrenal Hiperplazi Nedeni: Konjenital lipoid adrenal hiperplazi**
GÜRBÜZ F., TAŞTAN M., TURAN İ., YÜKSEL B., TOPALOĞLU A. K.
XXI. ULUSAL PEDİATRİK ENDOKRİNOLOJİ VE DİYABET KONGRESİ, Turkey, 26 - 30 April 2017

X. **CASR Mutasyonuna Bağlı Yenidoğanın Ciddi Hiperparatiroidisi'xxnde Cinacalset Tedavi Etkinliği**
TURAN İ., TAŞTAN M., kör y., KOTAN GEDİK L. D. , mert m. k. , GÜRBÜZ F., TOPALOĞLU A. K. , YÜKSEL B.
XXI. Ulusal Pediatrik Endokrinoloji ve Diyabet Kongresi, Turkey, 26 - 30 April 2017

XI. **Isolated Hypoaldosteronism: A Case Report**
TURAN İ., GÜRBÜZ F., TAŞTAN M., KOTAN GEDİK L. D. , TOPALOĞLU A. K. , YÜKSEL B.
2. Ege Endokrin Hastalıklar ve Genetik Sempozyumu, Turkey, 23 - 25 February 2017, vol.9

XII. **Inactivating Mutations in CCDC141 Causing Idiopathic Hypogonadotrophic Hypogonadism Kallmann Syndrome**
TURAN İ., hutchins i., hacıhamdioğlu b., özbek m. n. , KOTAN GEDİK L. D. , özkan y., stoner h., cheng p., GÜRBÜZ F., mengen e., et al.
espe 2016, 10 - 12 September 2016

XIII. **Idiopathic Hypogonadotrophic Hypogonadism Caused by Inactivating Mutations in SRA1**
KOTAN GEDİK L. D. , CARLTON C., DARCAN Ş., carr i., ÖZEN S., yan y., hamedani m., GÜRBÜZ F., mengen e., TURAN İ., et al.
espe 2016, 10 - 12 September 2016

## Citations

Total Citations (WOS):63
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