

# Prof. AYFER ALİKAŞİFOĞLU

## Personal Information

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## Education Information

Post Doctorate of Medicine, Hacettepe University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Çocuk Endokrinoloji Ve Metabolizma, Turkey 1993 - 1996

Expertise In Medicine, Hacettepe University, Tıp Fakültesi, Çocuk Sağlığı Ve Hastalıkları, Turkey 1988 - 1993

## Research Areas

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

## Academic Titles / Tasks

Professor, Hacettepe University, Hacettepe Tıp Fakültesi, 2007 - Continues

## Academic and Administrative Experience

Hacettepe Üniversitesi, Hacettepe Tıp Fakültesi, 2013 - 2014

## Published journal articles indexed by SCI, SSCI, and AHCI

- I. **Exercise performance in children and adolescents with cystic fibrosis with and without abnormal glucose tolerance: a single center cross-sectional study**  
Kocaaga E., Inal-Ince D., Dogru D., ALİKAŞİFOĞLU A., Ademhan-Tural D., Bozdemir-Ozel C., Calik-Kutukcu E., Saglam M., Vardar-Yagli N., Emiralioglu N.  
PHYSIOTHERAPY THEORY AND PRACTICE, vol.40, no.2, pp.230-240, 2024 (SCI-Expanded)
- II. **Alterations in insulin-like growth factor system in spinal muscular atrophy**  
YEŞBEK KAYMAZ A., Bal S. K., Bora G., TALİM B., Ozon A., ALİKAŞİFOĞLU A., Topaloglu H., YURTER H.  
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- III. **Feminizing Adrenocortical Tumors as a Rare Etiology of Isosexual/Contrasexual Pseudopuberty**  
VURALLI KARAOĞLAN D., Gonc N., Ozon A., EKİNCİ S., Dogan H. S., TEKGÜL S., ALİKAŞİFOĞLU A.  
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.14, no.1, pp.17-28, 2022 (SCI-Expanded)
- IV. **Management of prolactinomas in children and adolescents; which factors define the response to treatment?**  
ALİKAŞİFOĞLU A., ÇELİK ERTAŞ N. B., ÖZÖN Z. A., GÖNÇ E. N., KANDEMİR N.  
PITUITARY, vol.25, no.1, pp.167-179, 2022 (SCI-Expanded)
- V. **Sleep disordered breathing in patients with Prader willi syndrome: Impact of underlying genetic**

**mechanism**

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- VI. **Basal Serum Thyroxine Level should Guide Initial Thyroxine Replacement Dose in Neonates with Congenital Hypothyroidism**  
GÜNBEY C., Ozon A., GÖNÇ E. N., ALİKAŞİFOĞLU A., KARAHAN S., KANDEMİR N.  
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.13, no.3, pp.269-275, 2021 (SCI-Expanded)
- VII. **Cardiovascular risk factors in adolescents with type 1 diabetes: Prevalence and gender differences**  
VURALLI KARAOĞLAN D., Jalilova L., ALİKAŞİFOĞLU A., ÖZÖN Z. A., GÖNÇ E. N., KANDEMİR N.  
HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.236, 2021 (SCI-Expanded)
- VIII. **Epidemiology of type 1 diabetes in children and adolescents: a 50-year, single center experience**  
KANDEMİR N., VURALLI KARAOĞLAN D., ÖZÖN Z. A., GÖNÇ E. N., ARDIÇLI D., Jalilova L., GÜLÇEK Ö. N., ALİKAŞİFOĞLU A.  
HORMONE RESEARCH IN PAEDIATRICS, vol.94, no.SUPPL 1, pp.99, 2021 (SCI-Expanded)
- IX. **Clinical features and long-term follow up of childhood papillary thyroid cancer (PTC): a single reference-center experience**  
Ozon A., Emet D. C., Gonc N., VURALLI KARAOĞLAN D., Buyukyilmaz G., KANDEMİR N., ALİKAŞİFOĞLU A.  
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- X. **Is conventional treatment still the first choice in pediatric patients with PHEX mutations in an era of monoclonal FGF-23 antibody?**  
Alikasifoglu A., Unsal Y., Gonc N., Ozon A., Kandemir N., Alikasifoglu M.  
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- XI. **Long-term effect of conventional phosphate and calcitriol treatment on metabolic recovery and catch-up growth in children with PHEX mutation**  
ALİKAŞİFOĞLU A., ÜNSAL Y., GÖNÇ E. N., ÖZÖN Z. A., KANDEMİR N., ALİKAŞİFOĞLU M.  
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.34, no.12, pp.1573-1584, 2021 (SCI-Expanded)
- XII. **Approach to pheochromocytoma and paraganglioma in children and adolescents: A retrospective clinical study from a tertiary care center**  
ARDIÇLI B., USER İ. R., ÇİFTÇİ A. Ö., AKYÜZ C., KUTLUK M. T., Gonc N., ÖZÖN Z. A., ALİKAŞİFOĞLU A., OĞUZ B., HALILOĞLU M., et al.  
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- XIII. **Adrenocortical tumours in children: a review of surgical management at a tertiary care centre.**  
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ANZ journal of surgery, vol.91, no.5, pp.992-999, 2021 (SCI-Expanded)
- XIV. **Which parameters predict the beneficial effect of GnRHa treatment on height in girls with central precocious puberty?**  
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- XV. **Alpha-Melanocyte-Stimulating Hormone is Elevated in Hypothalamic Obesity Associated with Childhood Craniopharyngioma**  
Emet D. C., Ozon A., ALİKAŞİFOĞLU A., KANDEMİR N., Gonc N.  
OBESITY, vol.29, no.2, pp.402-408, 2021 (SCI-Expanded)
- XVI. **Obstructive sleep apnea in children with hypothalamic obesity: Evaluation of possible related factors**  
İYİGÜN İ., ALİKAŞİFOĞLU A., Gonc N., Ozon A., Eryilmaz Polat S., Hizal M., Kiper N., Ozcelik U.  
PEDIATRIC PULMONOLOGY, vol.55, no.12, pp.3532-3540, 2020 (SCI-Expanded)
- XVII. **Novel insights into diabetes mellitus due toDNAJC3-defect: Evolution of neurological and endocrine phenotype in the pediatric age group**  
ÖZÖN Z. A., ALİKAŞİFOĞLU A., KANDEMİR N., Aydin B., GÖNÇ E. N., KARAOĞLANOĞLU B., Celik N. B., Eroglu-

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- XVIII. **Gender-related differences in etiology of organic central precocious puberty**  
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- XIX. **Central nervous system imaging in girls with central precocious puberty: when is necessary?**  
VURALLI KARAOĞLAN D., GÖNÇ E. N., ALİKAŞİFOĞLU A., KANDEMİR N., ÖZÖN Z. A.  
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- XX. **Physical activity, functional capacity, and anaerobic power in cystic fibrosis with and without impaired glucose tolerance**  
Kocaaga E., Inal-Ince D., Ademhan-Tural D., Bozdemir-Ozel C., Calik-Kutukcu E., Vardar-Yagli N., SAĞLAM M., Dogru D., ALİKAŞİFOĞLU A., EMİRALİOĞLU N.  
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- XXI. **Oral health and halitosis among type 1 diabetic and healthy children**  
Iscan T. A., ÖZŞİN ÖZLER C., İLERİ KEÇELİ T., Guciz-Dogan B., ALİKAŞİFOĞLU A., UZAMIŞ TEKÇİÇEK M.  
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- XXII. **Treatment with Depot Leuprolide Acetate in Girls with Idiopathic Precocious Puberty: What Parameter should be Used in Deciding on the Initial Dose?**  
VURALLI KARAOĞLAN D., ALİKAŞİFOĞLU A., İYİĞÜN İ., Canoruc D., Ozon A., Gonc N., KANDEMİR N.  
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- XXIII. **Clinical and Molecular Analysis in 2 Families With Novel Compound Heterozygous SBP2 (SECISBP2) Mutations**  
Fu J., Korwutthikulrangsri M., GÖNÇ E. N., Sillers L., Liao X., ALİKAŞİFOĞLU A., KANDEMİR N., Menucci M. B., Burman K. D., Weiss R. E., et al.  
JOURNAL OF CLINICAL ENDOCRINOLOGY & METABOLISM, vol.105, no.3, 2020 (SCI-Expanded)
- XXIV. **Long-term effects of GnRH agonist treatment on body mass index in girls with idiopathic central precocious puberty**  
VURALLI KARAOĞLAN D., ÖZÖN Z. A., GÖNÇ E. N., ALİKAŞİFOĞLU A., KANDEMİR N.  
JOURNAL OF PEDIATRIC ENDOCRINOLOGY & METABOLISM, vol.33, no.1, pp.99-105, 2020 (SCI-Expanded)
- XXV. **Spondyloocular syndrome: Presentation of two siblings diagnosed with the rare disease and the results of Pamidronate Therapy**  
VURALLI KARAOĞLAN D., ŞİMŞEK KİPER P. Ö., Utine E., ÜNSAL Y., ALİKAŞİFOĞLU A., KANDEMİR N.  
HORMONE RESEARCH IN PAEDIATRICS, vol.91, pp.387, 2019 (SCI-Expanded)
- XXVI. **Improvement of final height in idiopathic central precocious puberty is associated with delay of bone maturation with GnRH agonist therapy under the age of 7 years**  
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- XXVII. **To whom should central nervous system imaging be performed in girls with central precocious puberty (CPP)?**  
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- XXVIII. **Long term effects of GnRH agonist therapy on BMI in girls with idiopathic central precocious puberty**  
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- XXIX. **Further expanding the mutational spectrum and investigation of genotype-phenotype correlation in 3M syndrome**  
Simsek-Kiper P. O., Taskiran E., KOŞUKCU C., ARSLAN U. E., Cormier-Daire V., Gonc N., Ozon A., ALİKAŞİFOĞLU A., KANDEMİR N., ÜTİNE G. E., et al.  
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- XXX. **Can having a sibling with type 1 diabetes cause disordered eating behaviors?**

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- XXXI. **What Is the Best Parameter to Decide the Initial Dose of Depot Leuprolide Acetate in Girls with Idiopathic Central Precocious Puberty?**  
VURALLI KARAOĞLAN D., ALİKAŞİFOĞLU A., Lyigun I., Canoruc D., Ozon A., Gonc N., KANDEMİR N.  
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- XXXII. **Gender-Related Differences in Etiological Distribution of Organic Causes of Central Precocious Puberty**  
VURALLI KARAOĞLAN D., Ozon A., Gonc N., KANDEMİR N., ALİKAŞİFOĞLU A.  
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- XXXIII. **Hyperinsulinemic Hypoglycemia in Congenital Disorder of Glycosylation Type-1a (CDG-1a)**  
Vuralli D., YILDIZ Y., SİVRİ H. S., ALİKAŞİFOĞLU A.  
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.362-363, 2018 (SCI-Expanded)
- XXXIV. **Early Onset GH Replacement in GH Deficiency: Is Neonatal Hypoglycemia Important for Long Term Follow-up?**  
ALİKAŞİFOĞLU A., CANORUÇ EMET S. D., Ozon A., Gonc N., KANDEMİR N.  
HORMONE RESEARCH IN PAEDIATRICS, vol.90, pp.386, 2018 (SCI-Expanded)
- XXXV. **The Frequency of Obstructive Sleep Apnea in Children with Hypothalamic and Exogenous Obesity**  
İYİGÜN İ., ALİKAŞİFOĞLU A., Ozon A., Gonc N., Hizal M., Eryilmaz S., Kiper N., Ozcelik U.  
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- XXXVI. **A Rare Cause of Hyperinsulinemic Hypoglycemia: Costello Syndrome**  
VURALLI KARAOĞLAN D., KOŞUKCU C., Taskiran E., Simsek P. O., ÜTİNE G. E., BODUROĞLU O. K., ALİKAŞİFOĞLU A., ALİKAŞİFOĞLU M.  
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- XXXVII. **Long-Term Follow-up of a Case with Proprotein Convertase 1/3 Deficiency: Transient Diabetes Mellitus with Intervening Diabetic Ketoacidosis During Growth Hormone Therapy**  
GÖNÇ E. N., Ozon A., ALİKAŞİFOĞLU A., KANDEMİR N.  
JOURNAL OF CLINICAL RESEARCH IN PEDIATRIC ENDOCRINOLOGY, vol.9, no.3, pp.283-287, 2017 (SCI-Expanded)
- XXXVIII. **Clinical and laboratory parameters predicting a requirement for the reevaluation of growth hormone status during growth hormone treatment Retesting early in the course of GH treatment**  
Vuralli D., GÖNÇ E. N., ÖZÖN Z. A., ALİKAŞİFOĞLU A., KANDEMİR N.  
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- XXXIX. **Clinical, genetic, and structural basis of congenital adrenal hyperplasia due to 11 beta-hydroxylase deficiency**  
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- XL. **A pheochromocytoma case diagnosed as adrenal incidentaloma**  
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- XLI. **Novel and prevalent CYP11B1 gene mutations in Turkish patients with 11-beta hydroxylase deficiency**  
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- XLII. **SOX9 GENE DUPLICATION-RELATED 46, XX OVOTESTICULAR DISORDER OF SEX DEVELOPMENT**  
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HORMONE RESEARCH IN PAEDIATRICS, vol.88, pp.371, 2017 (SCI-Expanded)
- XLIII. **ESTABLISHMENT OF HORMONE CONTROL WITH DEXAMETHASONE IN PATIENTS WITH CLASSICAL CONGENITAL ADRENAL HYPERPLASIA AFTER FINAL HEIGHT**  
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- XLIV. **HYPERCALCEMIA DEVELOPING IN THREE CASES FOLLOWING DISCONTINUATION OF DENOSUMAB TREATMENT**  
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- XLV. **Results of intraoperative gamma probe survey and frozen section in surgical treatment of parathyroid adenoma in children**  
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- XLVI. **Growth Hormone Deficiency in a Child with Neurofibromatosis-Noonan Syndrome.**  
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- XLVII. **An Adolescent Boy with Comorbid Anorexia Nervosa and Hashimoto Thyroiditis**  
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- XLVIII. **Development of Type 1 Diabetes in a Child with Inherited CD59 Deficiency Treated with Eculizumab**  
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- XLIX. **A Nonvirilized form of Classic 3 beta-Hydroxysteroid Dehydrogenase Deficiency Due to a Homozygous S218P Mutation in the HSD3B2 Gene in a Girl with Classic Phenylketonuria**  
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- L. **Hyperthyroidism After Allogeneic Hematopoietic Stem Cell Transplantation: A Report of Four Cases**  
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- LI. **Prevalence of nasal carriage of methicillin-resistant Staphylococcus aureus in children with diabetes mellitus: Trends between 2005 and 2013**  
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- LII. **Severe Undervirilisation in a 46,XY Case Due to a Novel Mutation in HSD17B3 Gene**  
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- LIII. **Evaluation of hypothalamic-pituitary function in children following acute bacterial meningitis**  
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- LIV. **Changing Etiological Trends in Male Precocious Puberty: Evaluation of 100 Cases with Central Precocious Puberty over the Last Decade**  
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- LV. **Variable Phenotype of Diabetes Mellitus in Siblings with a Homozygous PTF1A Enhancer Mutation**  
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- LVI. **Clinical characteristics of type 1 diabetes over a 40 year period in Turkey: secular trend towards earlier age of onset**  
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- LVII. **Need for Comprehensive Hormonal Workup in the Management of Adrenocortical Tumors in**

## Children

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- LIX. **Thyroid hormone resistance: a novel mutation in thyroid hormone receptor beta (THRB) gene - case report**  
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- LX. **Frequency of mutations in PROP-1 gene in Turkish children with combined pituitary hormone deficiency**  
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- XXXII. **Clinical and Molecular Analysis of 3M Syndrome In A Group of Turkish Patients**  
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- XXXIII. **Anorexia Nervosa ve Haşimoto Tiroiditi Beraberliği**  
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- ŞİMŞEK KİPER P. Ö., GÖNÇ E. N., KABAÇAM S., KARABULUT E., ÜTİNE G. E., COŞKUN T., ALİKAŞİFOĞLU A., HALİLOĞLU M., YILMAZ G., Project Supported by Higher Education Institutions, İskelet Displazilerinde Genetik Etiyolojinin Belirlenmesi, 2017 - 2019
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## Metrics

- Publication: 126  
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