

## Prof.Dr. SERAP DEMİRCİOĞLU

### Kişisel Bilgiler

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### Biyografi

Marmara Üniversitesi Çocuk Sağlığı ve Hastalıkları Bilim Dalında pediatri uzmanlık eğitimimi tamamladıktan sonra, aynı üniversitede Çocuk Endokrinoloji yan dal eğitimi aldım. Yan dal eğitimi sırasında uluslararası pek çok eğitim faaliyetine bilimsel toplantı ve bilim okulları olarak katılarak sunumlar yaptım. Yan dal eğitimimi tamamladıktan sonra klinik eğitimimi temel bilimler ile birleştirmek istedim ve de TÜBİTAK ve Fulbright'tan aldığım burslar ile Amerika Birleşik Devletleri Massachusetts General Hospital-Harvard Medical School'da hipofosfatemik rikets, psödohipoparatiroidi ve metilasyon bozuklukları alanında moleküler genetik çalışmalar yaptım. Türkiye'ye döndükten sonra bu konuda çalışmalarım ve farklı merkezler ile işbirliğim devam etti. 2010 yılında European Society for Paediatric Endocrinology (ESPE) 'Sabbatical Leave Programme' desteği ile 1 yıl süre yine aynı merkezde 'visiting scientist' olarak psödohipoparatiroidi (PHP) etiyo-patogenezi konusunda GNAS knock-out fare modellerinde çalışmalar yaptım. Halen, ESPE bünyesinde kurulan PHP grubunun aktif bir üyesi olarak Avrupa'da yapılan çalışmalara ve konsensus toplantılarına katılmaktayım. Avrupa Birliği Projesi olan COST Action BM10208- European Network for Human Congenital Imprinting Disorders çalışma grubunun aksiyon ortağı olarak Türkiye'yi temsil etmekteyim.

European Society for Paediatric Endocrinology (ESPE) tarafından Annual ESPE Meetingleri'nin program organizasyon komitesinde 4 yıl boyunca görev yapmak üzere 2010 yılında seçildim ve bu görevi 2014 yılında tamamladım. ESPE'nin bilim okulu olan Winter School'da 'Teaching Faculty' olarak 2015 yılında göreve başladım. European Society for Paediatric Endocrinology 1962 yılında Avrupa Pediatrik Endokrinoloji camiası tarafından kurulmuş olup, görev aldığım her iki pozisyon için de görev verilen ilk Türk bilim insanı olmam önem arz etmektedir.

İlk tip 1 diyabet prevalans çalışmasının ülkemizde okul çalışması kapsamında yapılmasında, sorumlu araştırmacı olarak görev aldım ve bu çalışmanın sonuçlarına göre okulda diyabet programının oluşturulmasında çalıştım. Diyabet 2020 projesi ve Sağlık Bakanlığı Diyabet programında raportör olarak görev aldım.

Journal of Pediatric Endocrinology and Metabolism ve Journal of Clinical Research in Pediatric Endocrinology dergilerinde yardımcı editörlük yapmanın yanı sıra pek çok dergi için yayın değerlendirmesi yapmaktayım. 2008'den bu yana Faculty of 1000 Prime'de Associate Faculty Member olarak yayın değerlendirmesi yapıyorum.

Ayrıca, Avrupa Birliği Çerçeve Programı 'imprinting Disorders' EURCOST çerçevesinde proje yürüttüm.

### Eğitim Bilgileri

Lisans Yandal, Marmara Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Türkiye 2001 - 2004

Lisans, Marmara Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, Türkiye 1995 - 2000

Ön Lisans, Hacettepe Üniversitesi, Tıp Fakültesi (İngilizce), Türkiye 1988 - 1995

### Yabancı Diller

İngilizce, A1 Başlangıç

## Araştırma Alanları

Sağlık Bilimleri

## Akademik Unvanlar / Görevler

Prof.Dr., Marmara Üniversitesi, Tıp Fakültesi, Dahili Tıp Bilimleri Bölümü, 2015 - Devam Ediyor

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## Yönetilen Tezler

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- XI. **Rare cause of severe hypertension in an adolescent boy presenting with short stature: Questions**  
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- XIV. **Low DHEAS Concentration in a Girl Presenting with Short Stature and Premature Pubarche: A Novel PAPSS2 Gene Mutation**  
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- XXVIII. **Age of obesity onset could be the first indicator of future metabolic complications - preliminary data of prospective multicenter study**  
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- XXX. **Genotype and Phenotype Characterization of Turkish Patients with Vitamin D Dependent Rickets Type IA**  
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- XXXI. **How to Approach Systemic Hypersensitivity reactions to Gonadotropin Releasing Hormone Analogues during treatment of Central Precocious Puberty**

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- XXXIII. **Rare Causes of Osteogenesis Imperfecta are Common in Consanguineous Pedigrees**  
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- XXXV. **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**  
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- XL. **Clinical and Laboratory Characteristics of Hyperprolactinemia in Children and Adolescents: National Survey**  
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- XLI. **PPP2R3C gene variants cause syndromic 46,XY gonadal dysgenesis and impaired spermatogenesis in humans**  
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- XLII. **Incidence of Type 1 Diabetes in Children Aged Below 18 Years during 2013-2015 in Northwest Turkey**  
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- XLIII. **THE CO-EXISTENCE OF TWO RARE DISEASES: A CASE REPORT**  
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**XLVII. The distribution of different types of diabetes in childhood: A single center experience**

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KAYGUSUZ S. B., Yesil G., KIRKGÖZ T., Turan S., BEREKET A., GÜRAN T.

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**L. An Unusual Cause of Short Stature**

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- XIX. **Disorders of sexual differentiation**  
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- XX. **Introduction to molecular endocrinology**  
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- LII. **hCG from an Occult Source Causing Peripheral Precocious Puberty Identification of the Tumour 6 Years After Presentation**  
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- LIII. **Neonatal Hypocalcemia**  
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- LIV. **From Pseudohypoparathyroidism to Inactivating PTH PTHrP Signaling Disorder iPPSD a Novel Classification Proposed by the European EuroPHP Network**  
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- LXVII. Disorder of Adrenocortical Excess and Adrenal Medulla**  
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