

# Öğr.Gör. FİLİZ BAŞAK ERGİN

## Kişisel Bilgiler

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## Eğitim Bilgileri

Doktora, Ankara Üniversitesi, Biyoteknoloji Enstitüsü, Temel Biyoteknoloji (Dr), Türkiye 2007 - 2010

Yüksek Lisans, Ankara Üniversitesi, Biyoteknoloji Enstitüsü, Temel Biyoteknoloji (YI) (Tezli), Türkiye 2004 - 2006

Lisans, Ankara Üniversitesi, Fen Fakültesi, Biyoloji Bölümü, Türkiye 1999 - 2004

## Yaptığı Tezler

Doktora, Sendromik olmayan otozomal resesif işitme kayıplı ve MYO15A genine bağlantı bulunan ailelerde mutasyon analizi, Ankara Üniversitesi, Biyoteknoloji Enstitüsü, Temel Biyoteknoloji (Dr), 2010

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## Araştırma Alanları

Sağlık Bilimleri

## Akademik Unvanlar / Görevler

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## SCI, SSCI ve AHCI İndekslerine Giren Dergilerde Yayınlanan Makaleler

- A possibly new autoinflammatory disease due to compound heterozygous phosphomevalonate kinase gene mutation**  
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## acidosis

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- V. **Hypophosphatasia: is it an underdiagnosed disease even by expert physicians?**  
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- XIV. **MPZL2 is a novel gene associated with autosomal recessive nonsyndromic moderate hearing loss**  
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- XVI. **Novel EYA1 variants causing Branchio-oto-renal syndrome**  
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- XXIV. **Identification of Copy Number Variants Through Whole-Exome Sequencing in Autosomal Recessive Nonsyndromic Hearing Loss**  
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- XL. **Analysis of NPHS2 mutations in Turkish steroid-resistant nephrotic syndrome patients**  
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## Diğer Dergilerde Yayınlanan Makaleler

- I. **A Turkish case of incontinentia pigmenti with a deletion mutation at Inhibitor of kappa B kinase gamma gene**  
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- II. **m.3010G>A Değişikliğinin Türk Populasyonunda Siklik Kusma Sendromuna Etkisi**  
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## Hakemli Kongre / Sempozyum Bildiri Kitaplarında Yer Alan Yayınlar

- I. **Ultra-Nadir Bir Hastalık ve Yeni Bir Mutasyon; Smg9 Eksikliği, Bir Aile 4Etkilenmiş Birey**  
Ergin F. B., İnci A., Ezgü F. S.  
6. Ulusal Çocuk Genetik Kongresini 09-12 Kasım 2023, Aydın, Türkiye, 9 - 11 Kasım 2023
- II. **Could Targeted Next Generation Sequencing Be A First Line Diagnostic Method for Lysosomal Storage Disease?**  
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- III. **Could Targeted Next Generation Sequencing Be A First Line Diagnostic Method for Lysosomal storage Diseases?**  
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## Metrikler

Yayın: 49

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