

PEDIATRIC GENETIC DISEASES



Assoc. Prof. Özlem Akgün Doğan, MD

Acibadem Altunizade Hospital, İstanbul · Acibadem Healthcare Group

MEDICALLY VERIFIED PROFILE



EXPERIENCE
20+ years



LANGUAGES
Turkish · English



CONSULTS
Video · On-site



PATIENTS
90+ countries



SPECIALTY
Pediatric Genetic Diseases



FOCUS
Achondroplasia · Rett syndrome



HOSPITAL
Acibadem Altunizade Hospital, İstanbul



DEPARTMENT
Pediatric Genetic Diseases



CLINICAL FOCUS

Dr. Doğan's clinical work centers on the evaluation and follow-up of children with genetic conditions. In achondroplasia, a genetic condition affecting bone growth, her role includes assessing the clinical features, discussing genetic testing where appropriate and helping families understand the diagnosis.

Because a genetic diagnosis can affect several aspects of a child's health, evaluation and follow-up may involve other pediatric specialties. Dr. Doğan discusses findings with families in a way that explains what a result does and does not mean, and outlines the options for further testing or follow-up.

Achondroplasia

Rett syndrome

Turner syndrome



CONDITIONS TREATED

Achondroplasia

Rett syndrome

Turner syndrome



EDUCATION & TRAINING

2017 Hacettepe University Faculty of Medicine Pediatric Genetic Diseases

2011 Dr. Sami Ulus Obstetrics and Gynecology, Child Health and Diseases Training and Research Hospital Child Health and...

2006 Hacettepe University Faculty of Medicine



PROFESSIONAL EXPERIENCE

2021 Acibadem Healthcare Services



PROCEDURES PERFORMED

ROP treatment



ACIBADEM HEALTHCARE GROUP Verified
network figures

45+

HOSPITALS & CLINICS

7

JCI-ACCREDITED

90+

COUNTRIES SERVED

20+

LANGUAGES IN-HOUSE



MEMBERSHIPS

- Pediatric Genetic Diseases Association
- European Society of Human Genetics



SELECTED PUBLICATIONS

- Comparison of the clinical diagnostic criteria and the results of the next-generation sequence gene panel in patients...
- H syndrome with a novel homozygous SLC29A3 mutation in two sisters. *Pediatr Dermatol.* 2020 Nov;37(6): 1135-1138.

10 publications listed on the online profile



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