



PEDIATRIC GENETIC DISEASES

Prof. Yasemin Alanay, MD

Acibadem Maslak Hospital, İstanbul · Acibadem Healthcare Group

MEDICALLY VERIFIED PROFILE

EXPERIENCE
30+ yearsLANGUAGES
Turkish · EnglishCONSULTS
Video · On-sitePATIENTS
90+ countries

SPECIALTY

Pediatric Genetic Diseases



FOCUS

Rare diseases ·
Hirschsprung disease

HOSPITAL

Acibadem Maslak Hospital,
İstanbul

DEPARTMENT

Pediatric Genetic Diseases



CLINICAL FOCUS

Dr. Alanay evaluates children with rare and inherited disorders. Her areas of expertise include skeletal dysplasias such as achondroplasia and osteogenesis imperfecta, as well as Ehlers–Danlos syndrome, Down syndrome, Hirschsprung disease, Harlequin fetus syndrome and Carney complex.

Genetic testing is a further part of her practice. This includes DNA testing, fetal DNA testing and the quadruple screening test. The choice of test, and whether testing is appropriate, depends on the individual clinical picture and family history and is discussed with the family during the evaluation.

Rare diseases

Hirschsprung disease

Harlequin fetus syndrome

Fetal DNA testing

Ehlers–Danlos syndrome



CONDITIONS TREATED

Rare diseases

Hirschsprung disease

Harlequin fetus syndrome

Ehlers–danlos syndrome

Down syndrome



EDUCATION & TRAINING

2011

Medical Specialization Board Pediatric Genetic Diseases

2008

Hacettepe University Faculty of Medicine — Associate Professor

2002

Hacettepe University Faculty of Medicine Pediatrics

1996

Hacettepe University Faculty of Medicine



PROFESSIONAL EXPERIENCE

2011

Acibadem Healthcare Services

2008–2011

Hacettepe University Faculty of Medicine, Pediatrics, Pediatric Genetics Unit, Associate Professor

2007–2008

Hacettepe University Faculty of Medicine, Pediatrics, Pediatric Genetics Unit, Assistant Associate Professor

2005–2006

Cedars–Sinai Medical Center–UCLA Intercampus Program, Research Fellowship in Skeletal Dysplasias

+4 earlier posts on the profile



PROCEDURES PERFORMED

Fetal DNA testing

Quadruple screening test

ROP treatment

ACIBADEM HEALTHCARE GROUP Verified
network figures

45+

HOSPITALS & CLINICS

7

JCI–ACCREDITED

90+

COUNTRIES SERVED

20+

LANGUAGES IN–HOUSE



MEMBERSHIPS

- International Skeletal Dysplasia Society
- European Society of Human Genetics
- Milli Pediatri Association
- Turkish Medical Association

+2 further societies on the profile



SELECTED PUBLICATIONS

- Genetic deficiency of tartrate-resistant acid phosphatase associated with skeletal dysplasia, cerebral calcifications...
- Molecular screening of ADAMTSL2 gene in 33 patients reveals the genetic heterogeneity of geleophysic dysplasia.

10 publications listed on the online profile



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+90 535 876 04 89

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acibademinternational.com

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FREE CONSULTATION

Ask Dr. Alanay about your case

Reply within 24 hours