MEDICAL GENETICS

Assoc. Prof. Ahmet Yeşilyurt, MD

Acibadem Maslak Hospital, İstanbul · Medical Genetics Department · Acibadem Healthcare Group

MEDICALLY VERIFIED PROFILE

EXPERIENCE	28+ years
LANGUAGES	Turkish · English
CONSULTS	Video · On-site
PATIENTS	90+ countries

SPECIALTY	FOCUS	HOSPITAL	DEPARTMENT
Medical Genetics	Genetic evaluation · Genetic diagnosis	Acibadem Maslak Hospital, İstanbul	Medical Genetics Department

CLINICAL FOCUS

Dr. Yeşilyurt's work covers genetic evaluation and genetic diagnosis for individuals and families, including prenatal genetic diagnosis and genetic testing on amniocentesis samples. His laboratory background includes tissue typing and transplant immunogenetic testing. He also evaluates chromosomal copy number abnormalities.

Among the inherited conditions listed in his areas of expertise are inherited thrombophilia, familial Mediterranean fever, maturity-onset diabetes of the young, Duchenne muscular dystrophy, nonclassical congenital adrenal hyperplasia, thyroid hormone resistance and hereditary glaucoma.

Genetic evaluation · Genetic diagnosis · Tissue typing · Transplant immunogenetic testing · Prenatal genetic diagnosis

CONDITIONS TREATED

Genetic evaluation · Genetic diagnosis · Tissue typing · Inherited thrombophilia · Familial mediterranean fever

EDUCATION & TRAINING

2007	Atatürk University Faculty of Medicine Medical Genetics
1998	Atatürk University Faculty of Medicine

PROFESSIONAL EXPERIENCE

2025	Acibadem Health Group
2019	Acibadem Maslak Hospital Medical Genetics Outpatient Clinic — Medical Genetics Specialist
2019	Acibadem Labgen Genetic Evaluation Center — Director
2019	Acibadem Labmed Ankara Tissue Typing Laboratory — Supervisor

+8 earlier posts on the profile

PROCEDURES PERFORMED

Transplant immunogenetic testing · Prenatal genetic diagnosis · Genetic testing (PGT/PGD)

ACIBADEM HEALTHCARE GROUP

Verified network figures

45+	7	90+	20+
HOSPITALS & CLINICS	JCI-ACCREDITED	COUNTRIES SERVED	LANGUAGES IN-HOUSE

MEMBERSHIPS

- American Society of Human Genetics
- Turkish Immunology Association
- Medical Genetics Association
- European Society of Human Genetics

+1 further societies on the profile

SELECTED PUBLICATIONS

- Homozygous SQSTM1 nonsense variant identified in a patient with brainstem involvement, Brain Dev.
- Association of vitamin D receptor gene FokI and TaqI polymorphisms and risk of RDS J Matern Fetal Neonatal Med.

30 publications listed on the online profile

PHONE	WHATSAPP	WEBSITE	FREE CONSULTATION
+90 535 876 04 89	+90 536 904 82 62	acibademinternational.com	Ask Dr. Yeşilyurt about your case
Tap to call	Tap to chat	Open the site	Reply within 24 hours