

Prof. Dr. Serdar Ceylaner is a medical doctor- a medical geneticist who focuses on rare and undiagnosed diseases for both diagnostic and scientific studies. He is the director, partner, and founder of Intergen Genetics and Rare Diseases Diagnosis and Research Center. Serdar Ceylaner is the Vice Chairman of the Rare and Undiagnosed Diseases Committee of the European Association of Medical Specialists (UEMS). He is a member of several other UEMS committees (Department of Medical Genetics, Medical Genetics Board Examination Committee, Multidisciplinary Adolescent Health Joint Committee).

He is the former president of the Medical Genetics Association of Turkey and a member of the board of directors between 2009 and 2017. Between 1997 and 2017, he was the founder of the Department of Genetics at Zekai Tahir Burak Women's Health and Training Hospital. He has focused on genetics and rare diseases and has studied in this field for 27 years. He established a center for diagnosis, research, and educational work in this field. Undiagnosed diseases, medical complications, and intensive care unit patients are his main research areas in recent years.

For nearly 30 years, I have been conducting studies in the fields of rare and undiagnosed diseases, genetic diseases, patient examination, genetic tests, genetic counseling, scientific research, R&D, education, and scientific policy. For 23 years, together with my wife Gülay Ceylaner, and the strong team of Intergen, under the logic of a scientific Institute, we have been performing studies that have affected the whole world.

It is our greatest happiness to be a part of a team with more than 500,000 patient experiences, identification of many new diseases, development of new materials and scientific methods, and experiences of more than a hundred projects and more than 350 scientific publications. Contributing to many international organizations such as the European Union of Medical Specialists-UEMS, the World Health Organization, and the International Organization for Undiagnosed Diseases-UDNI is one of our main aims. We have built our whole life in this direction and we are doing our best to pave the way for those who want to work in this field. Between 1997 and 2017, he was the founder of the Department of Genetics at Zekai Tahir Burak Women's Health and Training Hospital. He has focused on genetics and rare diseases and has studied in this field for 27 years. He established a center for diagnosis, research, and educational work in this field. Undiagnosed diseases, medical complications, and intensive care unit patients are his main research areas in recent years.



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Founder

