FELICIANO RAMOS

1983: M.D. Degree at the University of Extremadura Medical School (Spain)

1984-1988: Resident of Pediatrics. University Hospital, Zaragoza, Spain.

1985: National Award of Pediatric Investigation

1988: Ph.D. in Genetics at the University of Zaragoza Medical School, Zaragoza, Spain

1990-1992: Fulbright Scholarship. Postdoctoral Fellow. The Children´s Hospital of Philadelphia, Philadelphia, USA

1993-1996: Assistant Professor of Pediatrics. University of Zaragoza Medical School, Zaragoza, Spain

1995-present: Scientific advisor of the Spanish Association of families with Fragile X Syndrome

1996-2006: Full Professor of Pediatrics. University of Zaragoza Medical School, Zaragoza, Spain

2004-present: Member of the Advisory Board of the National Fragile X Foundation (USA)

2005-2013: President of the Spanish Society of Human Genetics

2006-present: Chair Professor of Pediatrics, University of Zaragoza Medical School, Zaragoza, Spain

2008-present: Member of the Experts Committee of the Spain´s National Strategy for Rare Diseases

2010-present: Coordinator of the Spain´s National Reference Center for ornelia de Lange Syndrome. Scientific advisor of the Spanish Association of families with Cornelia de Lange Syndrome

2011: Elected member of the Royal Academy of Medicine of Zaragoza

2012-present: Member of the Scientific Committee of the Biobanco de Aragón

2013- Coordinator of the Clinical Group associated to CIBERER at the Hospital Clínico Universitario “Lozano Blesa”

- Author of more than 100 papers, around half in SCI journals such as Nature enetics, American Journal of Human Genetics, Human Molecular Genetics, Journal of Medical Genetics or American Journal of Medical Genetics.

-Author of more than 40 chapters in Pediatrics and Genetics books.

-Principal Investigator of severalresearch projects funded by the Spain´s Ministry of Health (ISCIII-FIS) and Gobierno de Aragón.

-Member of the Spanish Pediatric Association (AEP), the American Society of Human Genetics (ASHG), the European Society of Human Genetics (ESHG) and the Spanish Association of Pediatrics (AEP).