



## Madelena Martin, M.D.

<b>Clinical Interests</b>	Dr. Madelena Martin's clinical interests are General Genetics and Biochemical Genetics.
<b>Research/Academic Interests</b>	Dr. Madelena Martin's research interests are General Genetics and Biochemical Genetics.
<b>Title</b>	Associate Professor, Department of Pediatrics
<b>Specialty</b>	General Genetics, Biochemical Genetics
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<b>Division</b>	Genomic Medicine
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<b>Education</b>	M.D., University of North Carolina at Chapel Hill School of Medicine, Chapel Hill NC 2000 B.A., California State University - Fresno, Fresno CA 1995
<b>Residency</b>	Genetics, Loma Linda University Medical Center, Loma Linda CA 2000-2003
<b>Fellowships</b>	Biochemical Genetics, UC San Francisco, San Francisco CA
<b>Board Certifications</b>	American Board of Medical Genetics - Biochemical Genetics American Board of Medical Genetics, Clinical Genetics American Board of Pediatrics
<b>Professional Memberships</b>	Society for Inherited Metabolic Disorders
<b>Honors and Awards</b>	Champions of Excellence, Pediatric Craniofacial Clinic Team, UMass Memorial, 2011 GlaxoSmithKline Endocrinology Scholar Award, Western Society of Ped Research, 2005 Second Year Resident of the Year, Pediatrics, Loma Linda University Med Center, 2002 Intern of the Year, Pediatrics, Loma Linda University Medical Center, 2001
<b>Select Recent Publications</b>	B Phung, A Suzuki, N Mans, J Welborn, M Martin. RET and PTEN: An Example in Understanding



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the Combinatorial Effects of Risk Variants in Cancer. (Platform) UCDMC 6th Annual Human Genomics Symposium. 2019 Nov.

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A Suzuki, J Sanine, S Collins, M Martin. 6.8Mb Xp deletion in a female with ornithine transcarbamylase deficiency pregnant with male fetus: Suggestions for pregnancy, delivery, and postnatal management. Society for Inherited Metabolic Disorders Annual Meeting, Seattle. 2019 Apr.

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Borzutzky A, Crompton B, Bergmann AK, Giliani S, Baxi S, Martin M, Neufeld EJ, Notarangelo LD. Reversible severe combined immunodeficiency phenotype secondary to a mutation of the proton-coupled folate transporter. *Clin Immunol*. 2009 Dec;133(3):287-94. doi:10.1016/j.clim.2009.08.006. Epub 2009 Sep 9. PMID:19740703.

Kronn D, Mofidi S, Braverman N, Harris K; Diagnostics Guidelines Work Group. Diagnostic guidelines for newborns who screen positive in newborn screening. *Genet Med*. 2010 Dec;12(12 Suppl):S251-5. doi:10.1097/GIM.0b013e3181fe5d8b. PMID:21150371.

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