Molecular and Medical Genetics Grand Rounds
2009 - 2010 Grand Rounds Schedule
Thursdays, 9:00 to 10:00 A.M., Mac Hall 3198

October 1  Jacob Reiss, M.D., Adjunct Associate Professor
“A Rare Variant Form of Ehlers-Danlos Syndrome”

October 8  Kory Keller, M.S., C.G.C., Genetic Counselor
“New Frontiers in NF1, NF2, & SPRED1”

October 15 Franklin Moore, M.D., Fellow, Pathology
“BRCA1: of Pathology, Patients, and Patents”

October 29 Chris Harrington, Ph.D., Sr. Staff Scientist
“DNA Microarrays: Past, Present, and Future”

November 5 Yassmine Akkari, Ph.D.
“The Cytogenetics of Multiple Myeloma”

November 12 Eric Orwoll, M.D., Associate Dean
“Genetic & Proteomic Studies of Osteoporosis and Sarcopenia”

November 19 Peter Francis, M.D., Associate Professor
“Treating Inherited Retinal Degenerations”

December 3 Cori Feist, M.S., C.G.C., Genetic Counselor
“TP63-related disorders”

December 10 Nicoleta Voian, M.D., Fellow, Genetics
“New Steps Toward Treatment of Duchenne Muscular Dystrophy – Exon Skipping Therapy”

December 17 Jessica Adsit, M.S., Professor
“Genetic and Epigenetic Implications of Assisted Reproductive Technology”

January 7  George Leonard, M.D., Fellow, Pathology
“Prenatal and Preimplantation Diagnosis of Hemoglobinopathies”

January 14 Kathryn Murray, M.S., Genetic Counselor
“Family History: Separating Truth from Action”

January 21 William Horton, M.D., Professor
“Progress in Understanding and Treating Achondroplasia”

January 28 Melanie Gillingham, Ph.D., Assistant Professor
“Feeding and Fasting in Alaska Native Children with CPT1A deficiency”

February 4  Stephen Moore, Ph.D., Assistant Professor
“Atypical Rett Syndrome”
February 11  Grover Bagby, M.D., Professor  
“Hematopoietic Dysfunction in Fanconi Anemia; Is it all about DNA damage?”

February 18  Cary Harding, M.D., Associate Professor  
“Disorders of Creative Biosynthesis & Transport”

March 4  Dana Kostiner, M.D., Assistant Professor  
“AAA Syndrome: It has nothing to do with roadside assistance”

March 11  Lynn Loriaux, M.D., Professor  
“Fuller Albright”

March 18  Dave Koeller, M.D., Professor  
“Newborn screening and treatment of Glutaric Acidemia Type 1”

April 1  Mary Williams, M.D., Adjunct Professor  
“The Ichthyoses: Pearls and Pitfalls for the Geneticist”

April 8  Gwen Fraley, M.S., Genetic Counselor  
“Perception of Risk: Discussing Autism Spectrum Disorders in the prenatal counseling setting”

April 15  Sue Richards, Ph.D., Professor  
“EGAPP/Evaluation of Genomic Applications in Practice and Prevention”

April 22  Nicoleta Voian, M.D., Fellow, Genetics  
“Prenatal Screening for Spinal Muscular Atrophy – A Comprehensive View”

April 29  Kelly Hamman, M.S., Genetic Counselor  
“Astrocytoma and Associated Inherited Cancer Syndromes”

May 6  Carrie Nielson, Ph.D, MPH, Assistant Professor  
“Genetic Epidemiology of Osteoporosis”

May 13  Monique Johnson Ph.D., Assistant Professor and  
Robert Searles, Ph.D, Director, Massively Parallel Sequencing Shared Resource  
“NextGen Sequencing: Will this innovative technology change the face of genetic research and clinical testing?”

May 20  William Noonan, M.D., J.D., Attorney, Klarquist Sparkman, LLP  
“Biological Patents, DNA Inventions, and Myriad Genetics”

May 27  Allison Gregory, M.S., Genetic Counselor  
Michael Kruer, M.D., Fellow  
“Update on Neurodegeneration with Brain Iron Accumulation: New Genes, New Directions”

June 3  Jon Zonana, M.D., Professor  
“Update of the Hypohidrotic Ectodermal Dysplasia (HED)”