



Photo: Jean-Marc Good

Genetic Medicine: developments and perspectives

Vendredi 1er décembre 13.00 - 19.00
CHUV Auditoire Jequier-Doge
entrée libre

**Symposium pour le départ de
Sheila Unger et Andrea Superti-Furga**

Ce petit symposium en anglais, organisé pour le départ de la Pr Sheila Unger et du Pr Andrea Superti-Furga, fait le point sur quelques aspects actuels de recherche et de pratique clinique en médecine génétique avec des orateurs prestigieux. Nous serons ravis de vous accueillir pour un moment scientifique et amical!

Christel Tran et François Clergue, Service de médecine génétique, CHUV

Programme

Manuel Pascual	Professor and Dean of Faculty; Faculty of Biology and Medicine, University of Lausanne	Welcome and Opening remarks
Sheila Unger	Former Associate Professor, University of Lausanne; Genetica AG, Lausanne	Introduction
Christine Hall	Emerita Professor of Radiology, Great Ormond Street Hospital, London, UK	The future of radiology, or: does radiology have a future?
Gen Nishimura	Former Director, Department of Radiology, Tokyo Metropolitan Children's Medical Center; Radiology Department, Musashino-Yowakai Hospital, Tokyo, Japan	A review of the history of bone dysplasias from a radiologist's viewpoint
Dan Cohn	Professor of Molecular, Cell, and Developmental Biology, and Orthopaedic Surgery; UCLA, Los Angeles, USA	Trespassing in the genome: Gene discovery in the skeletal dysplasias
Antonio Rossi	Professor of Biochemistry, Department of Molecular Medicine, University of Pavia, Italy	Intracellular sulfate – so simple, so important, so difficult to provide
Francesco Ramirez	Professor of Pharmacological Sciences, Orthopedics, Medicine and Cardiology; Icahn School of Medicine at Mount Sinai Hospital, New York, USA	TGFb signaling in Marfan syndrome: a tale of two stories : the long side
Valérie Cormier-Daire	Professeur de Genetique, Université Paris Descartes, INSERM, AP-HP ; Institut IMAGINE et Hôpital Necker-Enfants Malades, Paris, France	TGFbeta and Fibrillin pathways: the short side
Carlos Ferreira	Staff Clinician, Metabolic Medicine Branch, and Head, Skeletal Genomics Unit, NHGRI/NIH, Bethesda, USA	Genomics, models and therapeutic targets of genetic bone diseases
Livia Garavelli	Director, Medical Genetics Unit, Obstetrics and Pediatrics Department, AUSL IRCCS Reggio Emilia, Italy	The importance of networking in diagnosing rare disorders
*** Coffee break ***		
Marco Tartaglia	Head, Molecular Genetics and Functional Genomics, Ospedale pediatrico Bambino Gesù, Rome, Italy; and Dept. of Pediatrics, Mount Sinai School of Medicine, New York, USA	From Noonan syndrome to the Rasopathies – a long and winding pathway
Carlo Rivolta	Professor for Ophthalmic Genetics, University of Basel, CH; Professor of Medical Genetics, Department of Genetics and Genome Biology, University of Leicester, UK	Genetic bioinformatics from bench to bedside: a fruitful exchange
Matt Warman	Professor of Genetics, Harvard Medical School, and Department of Orthopedics, Boston Children's Hospital Precision Medicine Service, Boston, USA	Efficient in vivo testing for variants of uncertain significance (VUS)
Marc Tischkowitz	Department of Medical Genetics, University of Cambridge, UK	Polygenic – no longer Pollyanna?
Christel Tran	Privat-Dozent and Head of the Adult Metabolic Disease Unit, Division of Genetic Medicine, Lausanne University Hospital CHUV, Lausanne	Application of molecular therapies in genetic and metabolic disorders
Luisa Bonafé	Former Associate Professor, University of Lausanne; Head, Pediatric Medical Team, Regional Medical Services, National Invalidity Insurance, Vevey	Molecular treatments for rare genetic diseases: a challenge for the social insurance system
Marc Abramowicz	Professor and Head, Division of Genetic Medicine, HUG et UniGE, Geneva	Look back, look forward - the perspective of a neighbour and friend
Andrea Superti-Furga	Professor emeritus, University of Lausanne and Lausanne University Hospital; Genetica AG, Lausanne	Thanks and farewell

Crédit formation
SGMG: 6 heures

***** Apéritif *****