Rare Bone Disease TeleECHO Program

2021-2022



Physicians of all specialties, nurse practitioners, physician assistants, and other healthcare professionals with an interest in rare bone conditions are welcome to attend monthly case-based telementoring presentations and discussions. The series will focus on general and disease specific topics related to rare bone diseases.

Rare Bone Disease TeleECHO Program August 2021 - July 2022 First Thursday of each month; 3:00pm-4:30pm ET



In support of improving patient care, this activity has been planned and implemented by the Osteogenesis Imperfecta Foundation and Project ECHO®. Project ECHO® is jointly accredited by the Accreditation Council for Continuing Medical Education (ACCME), the Accreditation Council for Pharmacy Education (ACPE), and the American Nurses Credentialing Center (ANCC), to provide continuing education for the healthcare team

AMA Designation Statement

Project ECHO® designates this live activity for a maximum of 1.5 AMA PRA Category 1 Credit™. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

Interdisciplinary Faculty

Michael Collins, MD

Senior Investigator, Skeletal Disorders and Mineral Homeostasis Section, National Institute of Dental and Craniofacial Research/NIH

Michael Lewiecki, MD

Director, Bone Health TeleECHO, University of New Mexico Health Sciences Center

Frank Rauch, MD

Pediatrics, Shriners Hospital for Children – Canada; Professor of Pediatrics, McGill University Montreal, Qc, Canada

Eric. T. Rush, MD, FAAP, FACMG

Clinical Geneticist, Children's Mercy Kansas City; Associate Professor of Pediatrics, University of Missouri-Kansas City School of Medicine; Clinical Associate Professor of Medicine, University of Kansas School of Medicine

Jay Shapiro, MD

Consulting Physician, Department of Endocrinology and Diabetes, Walter Reed National Military Medical Center

Dolores Shoback, MD

San Francisco Dept of Veterans Affairs Medical Center, Professor of Medicine University of California, San Francisco

Laura Tosi, MD

TeleECHO Faculty Chair; Director, Bone Health Program, Children's National Hospital; Associate Professor of Orthopedics and Pediatrics, George Washington University

Michael Whyte, MD

Medical Director at the Center for Metabolic Bone Disease and Molecular Research at Shriners Hospitals for Children — St. Louis; Professor of Medicine, Pediatrics, and Genetics at Washington University School of Medicine

Presented by the OI Foundation in partnership with the Rare Bone Disease Alliance.

Register at www.oif.org/ECHO

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8/5/2021	Differential Diagnosis of Overgrowth Conditions Matthew Warman, MD Director, Orthopedic Research Laboratories Professor of Orthopedic Surgery Boston Children's Hospital
9/2/2021	Skeletal Presentations of Lyosomal Storage Diseases Patricia Dickson, MD Centennial Professor of Pediatrics and Genetics; Chief, Division of Genetics and Genomic Medicine, Washington University School of Medicine in St. Louis
10/14/2021	Paget's Disease of Bone Frederick R. Singer, MD <i>Director, Endocrine-Bone Disease Program St. Johns Cancer Institute</i> <i>Clinical Professor Medicine, UCLA</i>
11/4/2021	The Dysmorphology Exam for Skeletal Dysplasias Danita Velasco, MD, FAAP, FACMG Assistant Professor, Pediatrics and Genetics, Munroe- Meyer Institute for Genetics & Rehabilitation, University of Nebraska Medical Center
12/2/2021	Pulmonary Challenges in Ol Cathleen Raggio, MD Co-Director of the Kathryn O. and Alan C. Greenberg Center for Skeletal Dysplasias at Hospital for Special Surgery; Orthopedic Surgeon, Hospital for Special Surgery, New York, NY
1/6/2022	Genetic Testing – Emerging Diagnostic Technologies Emily G. Farrow, PhD, CGC Associate Professor of Pediatrics; UMKC School of Medicine; Director of Laboratory Operations Genomic Medicine Center, Children's Mercy Hospital
2/3/2022	Hypoparathyroidism Rachel Gafni, MD Senior Research Physician; Head, Mineral Homeostasis Unit, Skeletal Disorders and Mineral Homeostasis Section National Institutes of Dental and Craniofacial Research, NIH
3/3/2022	Dense Bone Diseases: Too Much Of A Bad Or Good Thing Michael Whyte, MD Medical Director at the Center for Metabolic Bone Disease and Molecular Research at Shriners Hospitals for Children — St. Louis; Professor of Medicine, Pediatrics, and Genetics at Washington University School of Medicine
4/7/2022	FOP and Anesthesia Zvi Grunwald, MD, FASA <i>Executive Director, The Jefferson Israel Center, The James D.</i> Wentzler Professor and Emeritus Chair of Anesthesiology; Professor of Pediatrics, Sidney Kimmel Medical College at Thomas Jefferson University
5/5/2022	Differential Diagnosis of non-XLH FGF 23 Disorders Sherri-Ann M. Burnett-Bowie, MD, MPH <i>Assistant Professor of Medicine, Harvard Medical</i> <i>School</i>
6/2/2022	Achondroplasia Michael B. Bober, MD, PhD Director, Skeletal Dysplasia Program, A.I. DuPont Hospital for Children; Professor of Pediatrics; Stanley Kimmel Medical College, Thomas Jefferson University
7/7/2022	MCTO Nina Ma, MD Ed and Jeannette Kerr Family Endowed Chair in Endocrinology; Director, Bone and Mineral Metabolism Program, Children's Hospital Colorado; Associate Professor of Pediatrics, University of Colorado School of Medicine

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