

Rare Bone Disease TeleECHO Program

2020-2021



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 2020 - July 2021

First Thursday of each month; 3:00pm-4:00pm ET



In support of improving patient care, 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 1.0 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

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

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

Register at www.oif.org/ECHO

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- 08/06/20** **Melorheostosis: The Genes Behind the Dripping Candle Wax**
Timothy Bhattachyrra, MD *Orthopaedic Surgery, NIAMS/NIH*
- 09/03/20** **Evaluation of Patients with Hyperphosphatemia**
Michael Collins, MD *Senior Investigator, Skeletal Disorders and Mineral Homeostasis Section, National Institute of Dental and Craniofacial Research/NIH*
- 10/01/20** **Mechanisms of Bone Loss in Complex Lymphatic Anomalies**
Michael Kelly, MD, PhD *Professor of Pediatrics, Northeast Ohio Medical University*
- 11/05/20** **Dental Concerns in Patients with Rare Bone Disorders**
Tim Wright, DDS, MS *Bawden Distinguished Professor, Division of Pediatrics and Public Health, Adams School of Dentistry, University of North Carolina*
- 12/03/20** **Generalized Arterial Calcification of Infancy (GACI)**
Carlos Ferreira, MD *Chief, Skeletal Genomics Unit, National Human Genome Research Institute, NIH*
- 01/07/21** **Skeletal Surveys – A Systematic Approach**
Dorothy Bulas, MD *Chief, Department of Radiology, Children’s National Hospital; Professor of Pediatrics and Radiology, George Washington University School of Medicine*
- 02/04/21** **Jansen’s Disease**
Harald Jueppner, MD *Chief, Pediatric Nephrology, Massachusetts General Hospital; Professor of Pediatrics, Harvard Medical School*
- 03/04/21** **Bone Pain in Children**
Alison Boyce, MD *Associate Research Physician, National Institute of Dental and Craniofacial Research, NIH*
- 04/01/21** **Multiple Hereditary Exostoses**
David S. Feldman, MD *Co-Director, Multiple Hereditary Exostoses Center, St. Mary’s Medical Center, West Palm Beach, FL*
- 05/06/21** **DXA Evaluation in the Child**
Catherine Gordon, MD *Adolescent Chief, Boston Children’s Hospital; Robert P. Masland, Jr. Chair of Adolescent Medicine, Professor of Pediatrics, Harvard Medical School*
- 06/03/21** **Adult Hypophosphatasia**
Kathryn Dahir, MD *Professor of Medicine, Vanderbilt University Medical Center*
- 07/01/21** **Evaluation of the Child with Multiple Fractures**
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*

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