

Project ECHO for Interstitial Lung Disease

Wednesday, December 21, 2022, 7am-8am

Medical Director	Robert Hallowell, MD
Principal Investigator	Aliaa Barakat, PhD
Didactic Speaker	Lida Hariri, MD
Session Facilitator	Robert Hallowell, MD
Target Audience	The series is designed for pulmonologists and radiologists, and for other health providers who care for people with ILD.

In accordance with HIPAA: All patient information will be de-identified during session.

Time	Didactic Presentation / Case Discussion	Presenter / Facilitator
7:00am-7:05am	Announcements and Introductions	Aliaa Barakat, PhD Robert Hallowell, MD
7:05am-7:25am	Didactic: Lung Biopsy in the Diagnosis of ILD	Lida Hariri, MD
7:25am-8:00am	Case Presentation and Discussion	Robert Hallowell, MD
8am	Closing Remarks	Arthur Dea, CCRP



In support of improving patient care, this activity has been planned and implemented by the Pulmonary Care and Research Collaborative, Ltd. 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.



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AMA Designation Statement

Project ECHO® designates this live activity for a maximum of 1.0 *AMA PRA Category 1 Credit™*. Physicians should claim only credit commensurate with the extent of their participation in the activity.

ECHO Activity ID: RSS330

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Disclosure Statement

Project ECHO®, in compliance with the ACCME Standards for Integrity and Independence in Accredited Continuing Education, requires that anyone who is in a position to control the content of an educational activity disclose all relevant financial relationships they have had within the last 24 months with an ineligible company.

None of the following planners / presenters for this educational activity have relevant financial relationship(s) to disclose with ineligible companies whose primary business is producing, marketing, selling, re-selling, or distributing healthcare products used by or on patients.

Arthur Dea, BA, CCRP; Fiona Gibbons, MD.

The following planners / presenters listed below have disclosed that they have a relevant financial relationship with an ineligible company.

Name	Nature of Relationship	Name of Ineligible Company
Aliaa Barakat, PhD	Principal Investigator of investigator-initiated research	Genentech, Inc.
Katharine Black, MD	Basic Science Researcher	Celgene/Bristol Myers Squibb, Bayer AG
Leo Ginns, MD	Site Principal Investigator for multicenter clinical trial	Nitto Denko
Robert Hallowell, MD	Site Principal Investigator or Co-Investigator for multicenter clinical trial	Galapagos NV, Hoffmann-La Roche, Nitto Denko
	Former Medical Advisory Board Member	Boehringer Ingelheim
	Editor and/or Content Contributor	Dynamed, UpToDate
Peter LaCamera, MD	Principal Investigator of investigator-initiated research	Boehringer Ingelheim
	Site Principal Investigator for multicenter clinical trial	Galapagos NV, Boehringer Ingelheim, Theravance Biopharma, Bristol Myers Squibb
Barry Shea, MD	Site Principal Investigator for multicenter clinical trial	Galapagos NV, FibroGen

All of the relevant financial relationships listed for these individuals have been mitigated.

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Name	Nature of Relationship	Name of Ineligible Company
Lida Hariri, MD, PhD	Principal Investigator of investigator-initiated research; Site PI for external collaborative research; Consultant preclinical and clinical therapy development	Boehringer Ingelheim
	Consultant preclinical and clinical therapy development	Pliant Therapeutics
	Consultant preclinical and clinical therapy development	Indalo Therapeutics
	Consultant preclinical and clinical therapy development	Bioclinica

All of the relevant financial relationships listed for these individuals have been mitigated.

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