Liver Forum Histology Webinar Series
Session 1
Best Practices: Increasing Reliability

Overview:
The level of discordance between pathologists, and the differences in operationalizing liver biopsy collection in trial protocols have been frequently raised as major issues and areas of concern for NASH clinical trials.

The goal of this session will be to discuss issues including: inter/intra-observer variability, scoring systems, and technical and logistical considerations.

Draft Agenda
Tuesday, September 1, 2020
12:00pm – 2:00pm ET

- Webinar Draft Agenda -		
Moderators: *Judith Ertle, Boehringer Ingelheim; Brent Tetri, Saint Louis University; Arun Sanyal, Virginia Commonwealth University*		
12:00 PM	**Introduction & Housekeeping**	*Katherine Barradas, Forum for Collaborative Research*
12:05 PM	**Setting the Stage: review of issues and recent data**	*Stephen Harrison, Oxford University*
12:20 PM	**Panel Discussion:**	*Pierre Bedossa, LiverPat*
		David Kleiner, NIH National Cancer Institute
		Oscar Cummings, Indiana University
		Cynthia Behling, University of California San Diego
		Massimo Siciliano, Università Cattolica del Sacro Cuore / external AIFA / EMA consultant
		Prakash Jha, U.S. Food and Drug Administration, CDRH
		Lara Dimick-Santos, U.S. Food and Drug Administration, CDER
1:20 PM	**Group Discussion**	*All*
2:00 PM	**Adjourn Webinar**	