

Biosimilars: A Decade of Experience and Future Directions Strategies for Improving Biosimilar Adoption and the Potential Role of Clinical Pharmacology

University of Maryland Center of Excellence in Regulatory Science and Innovation Food and Drug Administration Public Webinar April 13, 2022 | 1:00 – 3:30 pm Eastern Time

Agenda

Leading academic clinicians with specialties in oncology, rheumatology, gastroenterology, and endocrinology will share their experience with biosimilars, their perspectives on how to improve the efficiency of biosimilar evaluations, and how to increase biosimilar adoption, including the role of clinical pharmacology.

1:00 pm -	- 1:05 pm	Welcome
		Maureen Kane, PhD University of Maryland
1:05 pm -	– 1:10 pm	Opening Remarks
		Yow-Ming Wang, PhD Food and Drug Administration
1:10 pm -	- 1:35 pm	Biosimilars in Oncology
		Gary Lyman, MD, MPH, FACP, FRCP (Edin), FASCO University of Washington
1:35 pm -	- 2:00 pm	Biosimilars in Rheumatology (and Immune Mediated Inflammatory Diseases)
		Allan Gibofsky, MD, JD, MACR, FACP, FCLM Weill Cornell Medical College
2:00 pm -	– 2:25 pm	Biosimilars in IBD: Real World Evidence
		Gary Lichtenstein, MD, FACP, FACG, AGAF University of Pennsylvania
2:25 pm -	– 2:50 pm	Biosimilar Insulin Concepts
		Zachary Bloomgarden, MD, MACE, FASPC Icahn School of Medicine at Mount Sinai
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2:50 pm -	- 3:30 pm	Panel Discussion
	Moderator	Raymond Cross, MD, MS, AGAF, FACG University of Maryland
	Panelists	Yow-Ming Wang, PhD Food and Drug Administration
		Sarah Yim, MD Food and Drug Administration
		Zachary Bloomgarden, MD, MACE, FASPC Icahn School of Medicine at Mount Sinai
		Allan Gibofsky, MD, JD, MACR, FACP, FCLM Weill Cornell Medical College
		Gary Lichtenstein, MD, FACP, FACG, AGAF University of Pennsylvania
		Gary Lyman, MD, MPH, FACP, FRCP (Edin), FASCO University of Washington