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Biosimilars: The Next Frontier

Two-Part Webinar Series

August 12 & 13, 2020, 11:00 am – 2:00 pm PT

The launch of biosimilars in the US introduces competition to a trending specialty drug marketplace. Biosimilars are approved by the FDA for safety and are certified for "having no meaningful clinical difference" from their reference drugs. The nature of biologics makes it impossible to manufacture exact duplicates, as is the case with generics. Biosimilars is as much like a reference product as are distinct batches of the reference product! With savings estimated at \$54 billion¹ over ten years, we must address the market failures that are inhibiting adoption.

To accommodate current COVID-19 travel limitations, we have restructured this daylong symposium into two segments:

Welcome & Introduction to Biosimilar Webinar Series Kelly McDevitt					
2020	11:00-12:00 pm	Juliana Reed, MS, Vice President, Global Corporate Affairs Lead, Pfizer Wayne Winegarden, PhD, Sr Fellow & Director, Center for Medical Economics and innovation, Pacific Research Institute; Partner, Capital Economic Advisors	 Biosimilars: the big picture, the opportunity, the pipeline Overview of the landscape for biologics and biosimilars, key market trends and future issues for employers and manufacturers. Opportunities for biosimilar adoption and key factors to consider in the effective use of biosimilars for employers. Moderator: Kelly McDevitt 		
20	12:00	– 12:10 pm BREAK	Sponsor Slides/Acknowledgement		
August 12,	12:10-12:45 pm	Lucy Langer, MD Medical Director, Practice Solutions, The US Oncology Network	Biosimilars and Cancer Aligning incentives in Bundled Payment/CMS Oncology Care First Model Moderator: Anne Ladd		
Wednesday, August 12,	12:45-1:45 pm	Melissa Mantong, PharmD, MS, FCSHP, Pharmaceutical Consultant II, CalPERS Health Policy and Benefits Branch Clinical Policy and Programs Section Ladd Carman, BPharm, Director, Specialty Pharmacy Services, Costco Bill Dinger, MBA, CPA Director, Healthcare Innovation	 Benefit Design Strategies to Optimize Access and Affordability Purchasers will discuss how they have made decisions about the use of biosimilars in their pharmacy plan design. Key considerations in balancing the use of biosimilars and biologics. Moderator: Anne Ladd 		
		& Labor, The Walt Disney Company	1 A Louren Volo		
	1:45-2:00 pm Wrap-Up Day 1 Lauren Vela General Discussion and Q&A				

 $^{^{1} * \}texttt{https://www.rand.org/pubs/periodicals/health-quarterly/issues/v7/n4/03.html}$

Welcome to Day 2, Recap of Day 1 Lauren Vela					
	11:00-11:50 am	Kimberly Tsai, PharmD, CSP Clinical Pharmacist Supervisor, MemorialCare Medical Foundation Megan McIntyre, PharmD, MHA, Pharmacist Advisor, Benefits & Revenue Cycle, Virginia Mason Medical Center	 Provider Strategies to Manage Total Cost of Care Lessons from the field for purchasers. What are the issues with traditional cost plus reimbursement? How can purchasers help providers make high quality, cost-effective choices, including site of care? Moderator: Lauren Vela 		
0	11:5	50 am – 12:00 pm BREAK	Sponsor Slides/Acknowledgement		
Thursday, August 13, 2020	12:00-12:35 pm	Sameer Awsare, MD Associate Executive Director, The Permanente Medical Group	Kaiser's overall prescription drug management strategy. Case study of an evidence-based approach to adoption of biosimilars. Moderator: Kelly McDevitt		
	12:35-1:10 pm	Heather Sundar, PharmD, Executive Vice President of Medical Management at Archimedes, Inc.	Quantifying the Biosimilar Opportunity and Making Data-Driven Decisions How our intermediaries can help us and what the data tells us Moderator: Kelly McDevitt		
	1:10-1:45 pm	Brian Gifford, PhD, Director, Research and Analytics, Integrated Benefits Institute Rob Paczkowski, CEBS Senior Director, Global Benefits, eBay	 Impacting Workplace Productivity by Optimizing Affordability and Access Rheumatoid Arthritis: Opportunities and impact Measuring value by integrating the impact of adherence on lost worktime and lost productivity Moderator: Emma Hoo 		
		1:45-2:00 pm Wrap-Up Day	2 Kelly McDevitt		
		General Discussion and Q&A			

This program has been jointly developed by the Pacific Business Group on Health and Integrated Benefits Institute, with thanks to Program Sponsors

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