“The FDA and Cell-Based Therapies: Past, Present, and Future”

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Director, Center for Biologics Evaluation and Research
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Peter Marks, M.D., Ph.D. is the Director of the Center for Biologics Evaluation and Research (CBER) at the Food and Drug Administration. The center is responsible for assuring the safety and effectiveness of biological products, including vaccines, allergenic products, blood and blood products, and cellular, tissue, and gene therapies.

Peter Marks received his graduate degree in cell and molecular biology and his medical degree at New York University and completed Internal Medicine residency and Hematology/Medical Oncology training at Brigham and Women’s Hospital in Boston. He has worked in academic settings teaching and caring for patients and in industry on drug development. He joined the FDA in 2012 as Deputy Center Director for CBER and became Center Director in January 2016.

Tuesday, May 21, 2019 Clapp 108
4:30 – 5:30 p.m.

(This lecture is open to the University and its Affiliates)

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