

Monday, September 23

8:30 - 9:00 a.m.	Introduction and Welcome Remarks James C. Foster, <i>Chairman of the Board, President, and Chief Executive Officer, Charles River</i>
9:00 - 9:30 a.m.	2019 Award Winner: A Silicon Valley Approach to Understanding and Treating Disease Matt Wilsey, <i>Chairman, President, and Co-Founder, Grace Science Foundation</i>
9:30 - 10:15 a.m.	Keynote Session Brian Hubbard, PhD, <i>Chief Executive Officer, Dogma Therapeutics</i>
10:15 - 10:30 a.m.	Break
10:30 - 11:15 a.m.	Novel Approaches to Silence Disease Drivers Systemic Delivery of Investigational RNAi Therapeutics: Safety Considerations and Clinical Outcomes Peter Smith, PhD, <i>Senior Vice President, Early Development, Alnylam Pharmaceuticals</i>
11:15 a.m. - 12:00 p.m.	Novel Approaches to Silence Disease Drivers: Considerations for Viral Vector Manufacturing to Support Product Commercialization Richard Snyder, PhD, <i>Chief Scientific Officer and Founder, Brammer Bio</i>
12:00 - 1:00 p.m.	Lunch
1:00 - 1:45 p.m.	Accelerating Drug Discovery Through the Power of Microscopy Images Anne E. Carpenter, Ph.D., <i>Institute Scientist, Sr. Director, Imaging Platform, Merkin Institute Fellow, Broad Institute of Harvard and MIT</i>
1:45 - 2:30 p.m.	The Role of AI in Expediting Drug Discovery Target Identification for Nonalcoholic Steatohepatitis Using Machine Learning: The Case for nference Venky Soundararajan, PhD, <i>Founder and Chief Scientific Officer, nference/Qrativ</i>
2:30 - 2:45 p.m.	Break
2:45 - 3:30 p.m.	Technobite Sessions with Emulate Bio and University of Pittsburgh Drug Discovery Institute
3:30 - 4:15 p.m.	Artificial Intelligence Panel Discussion: Real World Applications from Discovery to Clinic Moderated by Carey Goldberg, <i>WBUR</i>
4:15 - 4:45 p.m.	Jack's Journey Jake and Elizabeth Burke, <i>Cure NF with Jack</i>
4:45 - 5:00 p.m.	Closing Remarks
5:00 - 6:00 p.m.	Networking Reception

Tuesday, September 24

8:45 - 9:00 a.m.	Opening Remarks and Recap James C. Foster, <i>Chairman of the Board, President, and Chief Executive Officer, Charles River</i>
9:00 - 9:30 a.m.	2018 Award Winner Update David Hysong, <i>Patient Founder and Chief Executive Officer, Shepherd Therapeutics</i> William Siders, <i>CDO, Shepherd Therapeutics</i>
9:30 - 10:15 a.m.	Advances in Human Genetics and Therapeutic Modalities Enable Novel Therapies Eric Green, <i>Vice President of Research and Development, Maze Therapeutics</i>
10:15 - 11:00 a.m.	How Genomics is Expediting Drug Discovery Manuel Rivas, <i>Assistant Professor, Department of Biomedical Data Science, Stanford University</i>
11:00 - 11:15 a.m.	Break
11:15 a.m. - 12:00 p.m.	Genomics Panel Discussion: Signposting Targets That Will Speed the Path to Market Moderated by Martin Mackay, <i>Co-Founder, RallyBio</i>
12:00 - 1:00 p.m.	Lunch
1:00 - 1:45 p.m.	Truly Personalized Medicines for Ultra-rare Diseases: New Opportunities in Genomic Medicine Timothy Yu, <i>Attending Physician, Division of Genetics and Genomics and Assistant Professor in Pediatrics, Boston Children's Hospital</i>
1:45 - 2:30 p.m.	Application of Machine Learning Technology for the Assessment of Bulbar Symptoms in ALS Fernando Vieira, <i>Chief Scientific Officer, ALS Therapy Development Institute</i>
2:30 - 2:45 p.m.	Break
2:45 - 3:30 p.m.	Accelerating Rare Disease Therapies Through the Regulatory Process Martine Zimmermann, <i>Senior Vice President and Head of Global Regulatory Affairs, Alexion Pharmaceuticals, Inc.</i>
3:30 - 4:00 p.m.	Wearing ALL the Hats: From Impossible to Possible Allyson Berent, <i>Chief Operating Officer, GeneTx Biotherapeutics</i>
4:00 - 4:15 p.m.	Closing Remarks