

FINAL AGENDA

Center Outcomes Forum

WebEx

Friday, November 20, 2020; 11:00 AM – 3:30 PM CDT

Time	Min	Topic	Speaker
11:00 – 11:10	10	Welcome and Introductions	J. Douglas Rizzo, MD, MS
11:10 – 12:10	60	Is minimal residual disease (MRD) for acute leukemia (AML, ALL) ready for use as a risk adjustment factor in the CSA? If not, what criteria need to be met in the future (e.g., consistent/standardized measures, etc.)? What changes in data collection should CIBMTR make to support future use of MRD?	Daniel Weisdorf, MD* Stella M. Davies, MBBS, PhD, MRCP^ Chris Hourigan, DM, DPhil^ Selina Luger, MD^ Wael Saber, MD, MS^ Bart Scott, MD^
12:10 – 12:20	10	Break	
12:20 – 1:20	60	Are there new approaches to account for social determinants of health in the CSA risk adjustment model?	Navneet Majhail, MD, MS* Kira Bona, MD, MPH^ Luciano Costa, MD, PhD^ Terri Hahn, PhD^ Richard Maziarz, MD^ Tatenda Mupfudze, PhD^ Ron Potts, MD^
1:20 - 1:40	20	Break	



Time	Min	Topic	Speaker
1:40 – 2:40	60	Can CIBMTR adequately adjust for the impact of the COVID-19 pandemic in the CSA? • What factors are most essential and how can they be incorporated in the risk adjustment?	John Wingard, MD*
			Kwang Woo Ang, PhD^ Christopher Dandoy, MD^
			Miguel Perales, MD [^]
			Bill Wood, MD, MPH^
2:40 – 2:50	10	Break	
2:50 – 3:10	20	CSA research project proposals (brief summary and plans)	Wael Saber, MD, MS
		□ Impact of CSA on HCT volumes	Akshay Sharma, MD
		□ Impact of CSA on HCT outcomes	Leslie Lehmann, MD
		 Impact of CSA on access/patient selection 	Mark Juckett, MD, MHCM
3:10 – 3:30	20	Update on consequences of CSA public reporting	Navneet Majhail, MD, MS
			J. Douglas Rizzo, MD, MS

^{*}Moderator ^Panelist