





Time: 13:00 - 14:00 CEST

Moderator: Rosa Giuliani, ESMO Director of Public Policy

Topic	Speaker	Organization	Time
Setting the scene	Rosa Giuliani	European Society for Medical Oncology (ESMO)	13:00 – 13:10
ICH guidelines and clinical research in Europe	Fergus Sweeney	European Medicines Agency (EMA)	13:10 – 13:20
Rationalising the bureaucratic burden in clinical trials	Jose Luis Perez-Gracia	University Clinic of Navarra (CUN)	13:20 – 13:30
Patient involvement in clinical research	Tanja Spanic	EUROPA DONNA & European Society for Medical Oncology (ESMO)	13:30 – 13:40
Case study: How can CROs facilitate non-commercial research?	Jeffrey Keefer	IQVIA	13:40 – 13:50
Discussion and Q&A	All		13:50 – 14:00
Conclusion and closing remarks	Rosa Giuliani	European Society for Medical Oncology (ESMO)	