European Epidemiological Forum

6 October 2020

16:15 - 16:30 **Conclusion**



The value of Real-World Evidence in Drug Life Cycle: promises, barriers and pathways to success

All times indicated on the Agenda are CEST

10:00 - 10:20	Welcome
10:20 - 10:45	Fostering trust among RWE stakeholders: Would medium to long-term partnerships help? Maurille Feudjo Tepie - Observational Research Director at Amgen
10:45 - 11:10	Critical thinking and Real world Evidence
	Elena Peruzzi - Evidence Generation & Data Analytics Head at Novartis Pharma
11:10 - 11:20	Break
11:20 - 11:45	NICE Guidance - Widening the evidence base: use of broader data
	and applied analytics in NICE's work
	Adrian Jonas - Associate Director for Data and Analytics at NICE
11:45 - 12:10	HTA, RWD and RWE: Validating and leveraging the different sources of evidence
	Oriol Solà-Morales - CEO at HITT- Health Innovation Technology Transfer
12:10 - 12:30	What did we learn this morning?
	Christian Agboton - Sr Global Brand Medical Director at Takeda
12:30 - 13:50	Lunch break
13:50 - 14:15	Value Based agreements & Innovative Contracting with Medicines:
	The value of Real-World Evidence for improving access & reimbursement
	Omar Ali - Visiting Lecturer Value Based Pricing at University of Portsmouth & Former Adviser to NICE
14:15 - 14:40	Data Transparency and Privacy in Real-World Research
	Raquel Billiones - Subject Matter Expert on Medical/Regulatory/Scientific Writing, Data Disclosure
	and Protection, Pharma/Medical Devices
14:40 - 15:05	Digitalization in Data Capturing Impacts Scalability of Patient Recruitment - Using
	the 2D Matrix Code of Outer Drug Packages as Patient Identifier for
	ePRO studies – The DePRO study
	Christian Müller - Teamleader Data Generation at Bayer
15:05 - 15:15	Break
15:15 - 16:15	WORKSHOP Covid-19 Pandemic: RWE, Opportunities and Challenges

