Thursday, September 21, 2023

Session 1	Ennov in 2022/2023 Overview – Sales & Marketing – Quality – Hosting & Security – Professionnal Services – Product Strategy				
	MORNING BREAK				
	Quality	Regulatory		Clinical	Human PV
Session 2	V10 Quality Core Model Presentation: Scope & Roadmap	Ennov Dossier v10.0	Samarind Product Update	Guided Tour to Ennov V10 EDC (1/2)	Ennov PV Human: what's new?
Session 3	Supplier Management: How to manage your supplier interactions and records in V10	Submission Document Core Model	Introduction to the Ennov platform for Samarind customers	Guided Tour to Ennov V10 EDC (2/2)	The Details of PV-Entry and PV-Admin
	AFTERNOON BREAK				
Session 4	V10 Document Management: How to manage your GMP documents trought the artifacts	V10 Ennov Analytics: Using and configuring KPIs for Regulatory	Ennov Clinical Product Update	MACRO Product Update	The Details of PV-Reporter

Friday, September 22, 2023

	Quality	Regulatory		Clinical	Human PV
Session 1	Quality Management: How to manage your deviations to CAPAs process	A quick taste of RIM data migration	Medical Devices & Combination products RIM Core Model	eTMF Completeness, Timeliness and Quality: From chaos to order	The Details of PV-Dashboard
	MORNING BREAK				
Session 2	V10 Learning Management: How to manage your training records	xEVMPD: Reg. affairs or Reg. ops?	Migrating from Samarind to Ennov Regulatory platform	Ennov CTMS: Driving insight and transparency in clinical trial management	Signal Detection and Signal Manager
	LUNCH BREAK				
Session 3	V10 Ennov Analytics: How to use your search and dashboard features	V10.x Regulatory wishlist		eLearning for Investigators and CRA	Integrating MACRO with Ennov CTMS	Management of Adverse Event in clinical trials. Reconciliation between the EDC and PV solutions.
	AFTERNOON BREAK				
Session 4	V10.x Quality Wishlist	IDMP and DADI 101 – be proactive!	Ennov Inbound / Outbound connector	V10.x Clinical wishlist	Other Topics: Reg Updates, Our Other PV Tools, Transitioning from PV Works