



INTERNATIONAL CONFERENCE OF DRUG REGULATORY AUTHORITIES

23-26 October 2012
Swissôtel, Tallinn, Estonia

ICDRA 23–26 October 2012 (regulators only)

	Tuesday / 23 October		Wednesday / 24 October	Thursday/ 25 October		Friday / 26 October
8.00–9.00	Registration					
9.00–10.30	PLENARY 1 Opening Ceremony		PLENARY 3 Ensuring Quality of Active Pharmaceutical Ingredients (+ including report from pre-ICDRA)	WORKSHOP G Assessing and responding to training needs of regulators	WORKSHOP H Responding to globalization of clinical trials	PLENARY 5 Pharmacovigilance: visions for future
Coffee						
11.00–12.30	PLENARY 2 Update on 14 th ICDRA recommendations and WHO progress report		PLENARY 4 Regulatory collaboration & networking	WORKSHOP I Regulatory harmonization	WORKSHOP J Patient and healthcare professional involvement in medicines regulation	PLENARY 6 Current topics
Lunch						
14.00–15.30	WORKSHOP A Current trends in regulating blood & cell therapies	WORKSHOP B Networking and collaboration for better regulation of herbal medicines	Keynote 14.00–14.50 City tours depart from Swissôtel 15.00	WORKSHOP K New tools for effective collaboration in combating SSFFCs	WORKSHOP L Should all regulators do everything? Best practices for prioritization and work-sharing.	PLENARY 7 Recommendations Closing remarks
	Coffee					
16.00–17.30	WORKSHOP C Collaboration and capacity building for vaccines regulation	WORKSHOP D Progress and challenges in regulating paediatric medicines		WORKSHOP M How should medical device products be regulated	WORKSHOP N Role of regulators in addressing availability of medicines	
18.00–20.00	ICDRA Welcome reception			Gala Dinner at 19.00 Estonia Concert Hall		