

Swiss Agency for Development and Cooperation SDC

INVITATION

A contribution to accelerating access to quality medical products in low and middle-income countries:

The Swissmedic Marketing Authorisation for Global Health Products (MAGHP) Procedure

Monday, 3 April, 14.30 – 16.30 (followed by a reception)
CICG, Centre International de Conférences Genève,
Rue de Varembé 17, 1211 Geneva



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Agenda

14.00 – 14.30	Coffee and Registration	
14.30 – 14.40	Welcome message	Sabrina Dallafior Matter
	The Swiss commitment to access to medical products	Ambassador, Deputy Permanent Representative of Switzerland to the UNOG and Permanent Representative of Switzerland to the Conference on Disarmament
14.40 –15.20	Setting the scene	
	Swissmedic Introduction: Swissmedic Procedure for Marketing Authorisation for Global Health Products (MAGHP)	Petra Doerr
		Deputy Director, Swissmedic, Swiss Agency for Therapeutic Products
	WHO Perspective: Overview of collaborative procedures and initiatives to register therapeutic products	Rutendo Kuwana
		Technical Officer, Regulatory System Strengthening Team, WHO
	Regulator's Perspective: Work sharing and reliance within the East African Community Medicines Regulatory Harmonization (EAC MRH) Programme and the cooperation with Swissmedic	Hiiti B. Sillo (tbc)
		Director General, Tanzania Food and Drugs Authority (TFDA)
15.20 – 16.20	Panel Discussion and Questions from the Audience	Alexander Schulze
	Christian Burri	(Moderation)
	Professor, Head Division Medicines Research, Department of Medicine Swiss Tropical & Public Health Institute	Co-Head, Global Programme Health Swiss Agency for Development and Cooperation - SDC
	Petra Doerr	
	Deputy Director, Swissmedic, Swiss Agency for Therapeutic Products	
	Murray M. Lumpkin	
	Deputy Director, Integrated Development (Regulatory Affairs), Lead for Global Regulatory Systems Initiatives, Bill & Melinda Gates Foundation	
	Hiiti B. Sillo (tbc)	
	Director General, Tanzania Food and Drugs Authority (TFDA)	
	Mike Ward	
	Coordinator, Regulatory System Strengthening Team, WHO	
	Rutendo Kuwana	
	Technical Officer, Regulatory System Strengthening Team, WHO	
16.20 – 16.30	Summary and Conclusion	Petra Doerr
		Deputy Director, Swissmedic, Swiss Agency for Therapeutic Products
16.30 – 18.00	Reception	