

Innovation

Affordability

Safety

Equity

Effective









Improving access to safe, effective and innovative quality medical devices



Research
Assessment

Training

Maintenance







Second WHO Global Forum

on Medical Devices:

Priority Medical Devices for Universal Health Coverage

Centre International de Conférences Genève (CICG) Geneva, Switzerland 22–24 November 2013



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Welcome

Distinguished ladies and gentlemen:

On behalf of the World Health Organization, we welcome you to Geneva, Switzerland for The Second WHO Global Forum on Medical Devices: Priority Medical Devices for Universal Health Coverage, providing the global public health community an opportunity for unprecedented collaboration to increase access to high-quality, safe, and appropriate-priority medical devices. Our global health initiative aims to impact all aspects of healthcare delivery, from diagnosis to treatment, and further enable those from academia, international organizations, industry, and NGOs to exchange their own experiences and challenges in our common endeavour to provide more equitable access to medical devices, particularly in limited-resource settings.

We thank you for your enthusiastic response to the call for abstracts, which has enabled us to organize a 3-day program that includes 28 workshops, 159 oral presentations in parallel sessions, 144 posters, and 8 films. You will join approximately 700 of your colleagues from 108 countries in 4 plenary sessions that will address the key issues facing the medical device community in 2013.

Thank you very kindly for your attendance and participation,

The Local Organizing Committee



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Meeting Objectives

- To define methods of increasing access to priority medical devices under the Universal Health Coverage initiative.
- To share evidence on best practices in health technology assessment, management and regulation of medical devices.
- To demonstrate the development and use of appropriate and innovative technologies that respond to global health priorities.
- To present the outcomes of the implementation of the World Health Assembly resolution on health technologies (WHA60.29) and the status of actions resulting from the First Global Forum on Medical Devices.

Background

The adoption of the first resolution on health technologies in May 2007 by the World Health Assembly (WHA 60.29) set the framework for an unprecedented focus on health technologies. Medical devices involve those health technologies that are critical to delivery within health systems. However, attention to issues of equity, quality and access is insufficient, and often the most essential medical devices are not available.

The 1st Global Forum on Medical Devices took place in Bangkok in September 2010, with participants coming from 107 Member States. The event raised awareness and served as a forum to share ideas on how to increase access to safe and effective medical devices.

Now, 3 years later, the 2nd Global Forum on Medical Devices will address the development of lists of medical devices by clinical intervention and disseminate information about innovative, appropriate, and affordable devices for low-resource settings in accordance with the WHA 60.29 resolution.

2nd WHO Global Forum on Medical Devices

22-24 NOVEMBER 2013 CICG, GENEVA, SWITZERLAND

Friday 22 November 2013

09:00—18:00	WORKSHOP TRACKS WHO/UN Tools to Improve Healthcare Delivery Health Technology Assessment Nomenclature, Standards and Regulations Health Tech Management and Clinical Engineering Radiation Safety / Biomedical Engineering	Salle 1 Salle 18 Salle 3 Salle 2
67.00 10.00	Health Tech Management and Clinical Engineering Radiation Safety / Biomedical Engineering Innovation	Salle 2 Salle 5/6 Salle 4
	Reproductive, Maternal, Newborn and Child Health / Approaches to Improving Healthcare Delivery	Salle 15

Saturday 23 November 2013

09:00-10:00	PLENARY SESSION 1 "Medical devices for Universal Health Coverage"		Salle 1
10:30-12:00	PARALLEL SESSIONS Health Technology Assessment: Networks and Socie Health Technology Management: Country Initiatives Regulation of Medical Devices Innovation in Medical Devices Health Care Delivery / Health Care Infrastructure		Salle 4 Salle 2 Salle 3 Salle 1 Salle 5/6
12:00-13:00	Lunch		
13:00-14:00	POSTER SESSION 1		
14:00—15:30	PARALLEL SESSIONS How to Prioritize Medical Devices Health Technology Management: Country Initiatives Regulation of Medical Devices: Country Initiatives Innovation in Medical Devices for Maternal and Chil	* Spanish translation available *	Salle 4 Salle 2 Salle 3 Salle 1 Salle 5/6
16:00-17:00	PLENARY SESSION 2 "The unifinished agenda: medical devices are indis	pensable for reaching the MDG targets"	Salle 1
17:00-19:00	Film viewing	Social networking / poster viewing	Salle 2

Sunday 24 November 2013

09:00-10:00	PLENARY SESSION 3 "Medical devices for the Non-Communicable Diseases (NCD) agenda"	Salle 1
10:30-12:00	PARALLEL SESSIONS Health Technology Assessment Policies for Medical Devices / Technical Specifications Safety of Medical Devices Innovative Medical Devices for Low-Resource Settings Human Resources in BME * French translation available *	Salle 4 Salle 2 Salle 3 Salle 1 Salle 5/6
12:00-13:00	Lunch	
13:00-14:00	POSTER SESSION 2	
14:00–15:30	PARALLEL SESSIONS Local Production in Low-Resource Settings Procurement of Medical Devices Patient safety / Medical Software Innovative Medical Devices for Low-Resource Settings Human Resources for Technology Life Cycle Management / Human Resources in Medical Physics	Salle 4 Salle 2 Salle 3 Salle 1 Salle 5/6
16:00–16:45	PLENARY SESSION 4 "Policies, innovation, regulation, assessment, management and safe use of medical devices for increasing access"	Salle 1
16:45-17:00	CLOSING SESSION Summary of recommendations from parallel sessions and closure of Forum	Salle 1
17:00	ADJOURN	

 $^{^{\}ast}$ All plenary sessions will have French and Spanish translation available *

WORKSHOP PROGRAMME (AS OF 18 NOVEMBER 2013) FRIDAY 22 NOVEMBER

09:00-09:10			PLENAI	RY / WELCOME / LOC	GISTICS		
	Salle 1	Salle 18	Salle 3	Salle 2	Salle 5/6	Salle 4	Salle 15
	WHO/UN Tools to Improve Healthcare Delivery	Health Technology Assessment	Nomenclature, Standards and Regulations	Health Tech Management / Clinical Engineering	Medical Imaging and Radiation Safety	Innovation	Reproductive, Maternal, Newborn and Child Health
09:15–10:00	Supporting integrated national strategic health planning, costing and health impact analysis: the OneHealth Tool (UNAIDS, UNDP, WHO, WB, UNFPA, UNICEF, the Futures Institute)	Creating synergies between national HTA and regional HTA agencies and hospitals in the assessment of medical devices (HTAi)	GMDN - a requirement for Unique Device Identification	Healthcare Technology Management (HTM): ACCE advanced clinical engineering	Role of medical physics in promoting radiation safety culture in health care	Innovation Sandbox Workshop:	MANDATE: priority setting for medical devices to reduce maternal, fetal and
10:00–10:45	Service Availability and Readiness Assessment (SARA) tool for health system planning and management (WHO)		(GMDN Agency)	workshops (ACCE)	(IOMP)	engaging medtech entrepreneurs to improve health in low- and middle- income countries through the power of co-creation (CAMTech,	neonatal mortality (RTI International)
10:45-11:10	Health break		Health break	Health break	Health break	Mass. General Hospital)	Health break
11:10–12:00	Crucial role of medical devices in emergency & essential surgical care (WHO)		Partnership on regulatory harmonization (AHWP, APEC)	Improving data quality and technology management with mobile devices (Health Partners International)	Medical imaging education in developing countries (ISR, WFUMB, ISRRT)	Hospital)	Medical device introduction: adding the Non-pneumatic Anti Shock Garment (NASG) for obstetric haemorrhage to programs and policies (UCSF)
12:00-13:30				Lunch / WHO visit			
	WHO/UN Tools to Improve Healthcare Delivery	Health Technology Assessment	Nomenclature, Standards and Regulations	Health Tech Management / Clinical Engineering	Biomedical Engineering	Innovation	Approaches to Improving Healthcare Delivery
13:30–14:20		for humanitarian How to set up an HTA	International standards — state	Computerized Maintenance	Human resources for		Améliorer les
	Kits for humanitarian health response	How to set up an HTA	of play and future trends in the medical domain (DITTA)	Management Systems (CMMS): essential features and pitfalls to avoid (ACCE)	medical devices: the role of the Biomedical Engineer (WHO)	Training for local innovation of affordable and appropriate medical devices in developing countries: learning	pratiques des projets d'appui à l'équipement médical intégrant des dons (Improving practices in medical equipment
14:30–15:20		How to set up an HTA agency (INAHTA)	trends in the medical domain	(CMMS): essential features and pitfalls to avoid	role of the Biomedical Engineer	innovation of affordable and appropriate medical	projets d'appui à l'équipement médical intégrant des dons (Improving practices
14:30–15:20 15:20–16:00	health response (WHO/UNFPA/	agency	trends in the medical domain (DITTA) Medical software — regulatory and legal trends	(CMMS): essential features and pitfalls to avoid (ACCE) WHO template for technical specifications of medical equipment	role of the Biomedical Engineer (WHO) How to define the basic academic curriculum to train clinical engineers	innovation of affordable and appropriate medical devices in developing countries: learning from the Stanford India Biodesign Experience (All India Inst. of	projets d'appui à l'équipement médical intégrant des dons (Improving practices in medical equipment support projects which include donations) (HUMATEM) * in French and English,
	health response (WHO/UNFPA/	agency	trends in the medical domain (DITTA) Medical software — regulatory and legal trends	(CMMS): essential features and pitfalls to avoid (ACCE) WHO template for technical specifications of medical equipment (WHO)	role of the Biomedical Engineer (WHO) How to define the basic academic curriculum to train clinical engineers	innovation of affordable and appropriate medical devices in developing countries: learning from the Stanford India Biodesign Experience (All India Inst. of	projets d'appui à l'équipement médical intégrant des dons (Improving practices in medical equipment support projects which include donations) (HUMATEM) * in French and English,

Useful information

Forum venue

Centre International de Conférences Genève Rue de Varembé 17 1211 Geneva

Internet

WiFi login: OMS Password: WHO



For SpotMe updates

Username: WHO13 Password: geneva13

Transportation to the CICG: Near all pre-booked hotels (except Ibis Nations), tram 15 can take you in the direction of the CICG terminus stop *Nations*. From Ibis Nations, bus 8 can transport you to the CICG stop *UIT* in the direction OMS/Appia.

Food at the venue

CICG restaurant in the venue offers reasonably priced meals (around 10-20 CHF).

Currency

Local currency is Swiss Francs (CHF) not Euros. Exchange rate is around US \$ 0.93.

Geneva tourism sites:

http://www.geneve-tourisme.ch http://myswitzerland.com http://www.geneva.com http://search.ch.com

Geneva transportation:

Geneva bus and tram information: http://www.tpg.ch/

Swiss train information: http://www.sbb.ch/ticketshop

Taxi stands are at almost all main squares. The Forum venue (CICG) is about 10 minutes from the airport (25-30 CHF). A taxi from the airport to the town centre costs 35–50 CHF.

Taxi-phone SA Geneva: +41 22 33 141 33 www.taxi-phone.ch AA Genève Central Taxi: +41 22 3 202 202 www.geneve-taxi.ch

Presenter guidelines

Speakers

- Speakers should deliver their slides to conference organizers the day before the presentation or before the start of the plenary session the morning of their talk.
- Speakers should arrive 15 minutes prior to the session start.
- To allow sufficient time for panel discussion, time limits will be strictly enforced.
- Speakers are expected to be present for the duration of the session.

Posters

- Posters should be set up before 08:00 on Saturday 23 November.
- Posters must be removed between 17:00 and 19:00 on 24 November.
- Poster presenters should be at the designated poster area at least 10 minutes before their assigned poster session begins and, if possible, during the lunch hour and coffee breaks.

Films

- Film presenters should arrive 15 minutes prior to the start of the film viewing.
- Each presenter may give a brief (1 minute or less) introduction before his/her video.
- After all films are completed, film presenters will participate in panel discussion.

Collaborating organizations

UN agencies







Non-governmental organizations in official relations with WHO

International Federation for Medical and Biological Engineering



International Federation of Hospital Engineering



International Society of Radiology



International Society of Radiographers & Radiological Technologists



International Union of Architects



World Federation for Ultrasound in Medicine and Biology



Organizations with memoranda of understanding with WHO

International Information Network on New and Emerging Health Technologies



Health Technology Assessment International



International Network of Agencies for Health Technology Assessment



Organization of Medical Physics



Other collaborating organizations

American College of Clinical Engineering



Diagnostic Imaging, Healthcare IT & Radiation Therapy Trade Association



Global Medical Technology Alliance





Institute of Biomedical Technology



International Union for Physical and Engineering Sciences in Medicine





