

PROMOTING EQUITABLE ACCESS TO HEALTH PRODUCTS THROUGH INNOVATION AND COLLABORATION

“A NEXUS FOR SUSTAINABLE UNIVERSAL HEALTH COVERAGE”

Joint UNICEF, UNFPA and WHO meeting with manufacturers and suppliers of
in vitro diagnostic products, vaccines & immunization devices, finished pharmaceutical products, active
pharmaceutical ingredients, contraceptive devices and vector control products

The 27 November-01 December 2023 Joint Meeting will be a face-to-face meeting taking place at
UN City, Marmorvej 51, 2100 Copenhagen, Denmark

DRAFT AGENDA

DAY 1	MONDAY, 27 NOVEMBER 2023
07:00–08:30	MEETING REGISTRATION: ENTRANCE TO UN CITY
SESSION 1 - PLENARY: 08:45–10:30 AUDITORIUM I, II & III	ADVANCING EQUITABLE ACCESS TO QUALITY HEALTH PRODUCTS THROUGH INNOVATIONS AND COLLABORATIONS <i>Facilitator: Marta Seoane, WHO</i> <i>Hybrid</i>
08:45–09:00	Administrative / Security arrangements <i>Henrik Pantmann, Head of Security – UN City</i>
09:00–09:30	Welcome from meeting host agencies: <i>WHO - UNICEF – UNFPA</i> <ul style="list-style-type: none"> • <i>Yukiko Nakatani, ADG/MHP</i> • <i>Karen Hong, Chief Supply Chain Management Unit, UNFPA</i> • <i>Leila Pakkala Director, UNICEF Supply Division</i>
09:30–09:45	Outline of the meeting Objectives, Agenda, Themes and expected outcomes <i>Deus Mubangizi, Unit Head, PQT, WHO</i>
09:45–10:00	Remarks and introduction of Keynote speaker <i>Marta Seoane, WHO</i>
10:00–10:30	Keynote address <i>Sir Jeremy Farrar, Chief Scientist, World Health Organization</i>
10:30–11:00	Coffee /Tea break

SESSION 2 - PLENARY: 11:00–13:00 AUDITORIUM I, II & III	BREAKING BARRIERS: ENSURING AFFORDABLE AND EQUITABLE ACCESS TO INNOVATIVE AND QUALITY ASSURED HEALTH PRODUCTS GLOBALLY <i>Facilitator: Marta Seoane, WHO</i> <i>Hybrid</i>
11:00-11:05	Introduction- <i>Deus Mubangizi, Unit Head, PQT, WHO</i>
11:05–12:15	Streamlining WHO Processes to facilitate new product innovation (-30 min) Speaker(s): Yukiko Nakatani, <i>ADG/MHP</i> and Jerome Salomon, <i>ADG/ULC</i> Moderated by Mubashar Update and progress on the Coordinated Scientific Advice (CSA)- Mercedes (-10 min) Presentations on new product pipelines (-30 minutes) HIV (DOHERTY, Meg <dohertym@who.int) TB (KASAEVA, Tereza <kasaevat@who.int) Malaria (NGAMIJE, Madandi Daniel <ngamijed@who.int) AMR (DR. GETAHUN, Haileyesus/DR. VAN WEEZENBEEK, Catharina)
12:15–12:45	Accelerating access to medicines for children <ul style="list-style-type: none"> • Welcome and framing <i>Martina Penazzato (WHO)</i> • Update on PADO TB deliberations <i>Tiziana Masini (WHO)</i> Panel Discussion: Accelerating access to medicines for children <ul style="list-style-type: none"> • <i>Sandra Nobre (MPP)</i> • <i>Jennifer Cohen (GARD-P)</i> • <i>Luwei Pearson (Unicef)</i> • <i>Fabienne Benoist (Dndi)</i> • <i>Cipla</i> Wrap up
12:45-13:00	Q&A
13:00–14:00	Lunch
SESSION 3 - PLENARY 14:00–15:30 AUDITORIUM I, II & III	INNOVATIVE PARTNERSHIPS IN LOCAL PRODUCTION, PROCUREMENT, AND REGULATORY ARRANGEMENTS <i>Facilitator: Marta Seoane, WHO</i> <i>Hybrid</i>
14:00-14:05	Introduction <i>Deus Mubangizi, Unit Head, PQT, WHO</i>
14:05-14:20	Outcomes from World Local Production Conference <i>Jicui Dong, Unit Head, WHO</i>
14:20–14:45	UNICEF roadmap on local production and local procurement <ul style="list-style-type: none"> • <i>Sebastian Meaney, Chief, Markets and Supply Financing</i> • <i>Hamadou Dicko, Snr Advisor, Supply,, AU Hub</i>
14.45-15.10	Global Fund-PEPFAR-UNITAID Initiative on HIV Diagnostics produced in Africa <i>Hui Yang, Global Fund</i>
15:10–15:30	Questions & Comments

15:30–16:00	Coffee /Tea break
SESSION 4 – PLENARY 16:00– 17:00 AUDITORIUM I, II & III	ACCELERATING HEALTHY SUPPLY CHAIN INTERVENTIONS AT GLOBAL, REGIONAL AND COUNTRY LEVEL <i>Facilitator: Marta Seoane, WHO and Deus Mubangizi, Unit Head, PQT, WHO</i> <i>Hybrid</i>
16:00–17:00	<ul style="list-style-type: none"> Managing commodity shortages and accessing information: Database on shortages - <i>Lisa Headman (WHO)</i> Improving sustainable access to antibiotics through market intelligence and coordination - <i>Jennifer Cohn (GARDP)</i> UNFPA's approach to quantification and collaborative national supply planning-<i>Stephen Mawa, Team Lead, Demand and Supply Planning, UNFPA</i> The Power of Traceability and Verification - <i>Max Kabalisa, UNICEF</i>
17:00–17:30	Questions & Comments
17:30–18:00 AUDITORIUM I, II & III	WRAP-UP OF DAY 1 AND PLANS FOR DAY 2 <ul style="list-style-type: none"> <i>Facilitator: Marta Seoane, WHO</i> <i>Deus Mubangizi - Unit Head, PQT, WHO</i>

DAY 2	TUESDAY, 28 NOVEMBER 2022
07:00–08:40	ENTRANCE TO UN CITY
SESSION 5 08:45–13:00	WHO AND UNFPA PREQUALIFICATION UPDATES-PARALLEL SESSIONS
SESSION 5.1 – PARALLEL TRACK 08:45–13:00	WHO IN VITRO DIAGNOSTICS (IVDs) PREQUALIFICATION TRACK <i>Session chair: Irena Prat, Team Lead, PQT/IVD, WHO</i> <i>Attendance mode: Hybrid</i>
Updates on Prequalification of in vitro diagnostics	
08:45–11:00	<p>Introduction & Welcome <i>Irena Prat, PQT/IVD, WHO</i></p> <p>Product dossier <i>Speaker: Mark Lanigan, PQT/IVD, WHO</i></p> <p>Performance evaluations <i>Speaker: Anne-Laure Page, PQT/IVD, WHO</i></p> <p>Labelling review and Public Reports <i>Speaker: Charles Chiku, PQT/IVD, WHO</i></p> <p>ePQS <i>Speaker: Helena Ardura, PQT/IVD, WHO</i></p> <p>Q&A</p>
11:00–11:30	Coffee/Tea break

11:30 – 13:00	<p>PQ Technical specifications <i>Speaker: Ute Ströher, PQT/IVD, WHO</i></p> <p>Change requests <i>Speakers: Fatima Gruszka and Helena Ardura, PQT/IVD, WHO</i></p> <p>Collaborative registration procedure for IVDs <i>Speaker: Susie Braniff, PQT/IVD, WHO</i></p> <p>Q&A</p> <p>Wrap Up <i>Irena Prat, PQT/IVD, WHO</i></p>
13:00–14:00	Lunch
SESSION 5.2 – PARALLEL TRACK 08:45–13:00	WHO MEDICINES PREQUALIFICATION TRACK <i>Session Chair: Matthias Stahl, Team Lead, PQT/MED, WHO</i> <i>Attendance mode: In-person only</i>
08:45–09:00	Introduction <i>Speaker: Matthias Stahl, Team Lead, PQT/MED, WHO</i>
09:00–09:25	Quality (Finished Pharmaceutical Products) <i>Speaker: Wondiyfraw Worku, PQT/MED, WHO</i>
09:25–09:50	Quality (Active Pharmaceutical Ingredients) <i>Speaker: Antony Fake, PQT/MED, WHO</i>
09:50–10:15	Bioequivalence <i>Speaker: John Gordon, PQT/MED, WHO</i>
10:15–10:40	Biotherapeutics & Biosimilars <i>Speaker: Guido Pante, PQT/MED, WHO</i>
10:40–11:05	WHO Public Assessment Reports (WHOPARs) <i>Speaker: Regine Lehnert, PQT/MED, WHO</i>
11:05–11:35	Q&A
13:00–14:00	Lunch
SESSION 5.3 – PARALLEL TRACK 08:45–13:00	WHO VACCINES & IMMUNIZATION DEVICES PREQUALIFICATION TRACK <i>Session Chair: Carmen Rodriguez Hernandez, Team Lead, PQT/VAX, WHO</i> <i>Attendance mode: Hybrid</i>
08:45–09:00	WHO vaccines prequalification overview
09:00–10:20	Vaccine assessment: CMC (chemistry, manufacturing and control), quality and post PQ activities and clinical data <i>Speakers: Olivier Lapujade, Rolando Dominguez PQT/VAX, WHO</i>
10:20–10:40	Programmatic suitability for PQ <i>Speaker: Emma Hernandez, PQT/VAX, WHO</i>
10:40–11:10	Facilitated discussions

11:10–11:30	Coffee/Tea break
11:30–12:00	Risk benefit assessment procedures. Updates on EUL <i>Speaker: Carmen Rodriguez Hernandez, Team Lead, PQT/VAX, WHO</i>
12:00–12:20	ePQS and eCTD updates applicable for VAX <i>Speaker: Emma Hernandez, PQT/VAX, WHO</i>
12:20–13:00	Facilitated discussions
13:00–14:00	Lunch
SESSION 5.4 – PARALLEL TRACK 08:45 – 13:00	WHO VECTOR CONTROL PRODUCT PREQUALIFICATION TRACK <i>Session Chair: A.g Team Lead, Dominic Schuler PQT/VCP, WHO</i> <i>Attendance mode: Hybrid</i>
08:45–09:00	<i>PQT/VCP – General Updates and Workplan for 2024</i>
09:00–11:00	<i>Procedures related to the evaluation of technical materials through JMPS and acceptance of source materials for use in formulation of VCPs</i>
11:00–11:30	Coffee/Tea break
11:30–12:30	<i>Presentation on the WHO Guideline for the Prequalification of ITNs and associated implementation documents</i>
12:30–13:00	<i>Questions and Answers</i>
13:00–14:00	Lunch
SESSION 5.5 – PARALLEL TRACK 08:45 – 15:00	UNFPA PREQUALIFICATION OF CONTRACEPTIVE DEVICES & MARKETS TRACK <i>Session Chair: Linda Serwaa, Head, Quality Assurance, Technical and WHO/UNFPA Prequalification, UNFPA</i> <i>Attendance mode: Hybrid</i>
08:45-09:05	WHO/UNFPA Prequalification (PQ) program updates <i>Speaker: Ashley Moyo, Technical Analyst, UNFPA</i>
09:05–09:40	summary of analysis of quality complaints <i>Speaker: David Hill, technical Expert for the UNFPA PQ Programme</i>
09:40–10:00	summary of analysis of reported significant changes <i>Speaker: David Hill, technical Expert for the UNFPA PQ Programme</i>
10:00–10:30	Post-Market Surveillance Programme for PQ'ed manufacturers <i>Speaker: Dr. Kamakshinatha Sivakumar, technical Expert for the UNFPA PQ Programme</i>
10:30–11:00	Technical support for new PQ applications – overview/outline, eligibility, implementation etc., <i>Speaker: Dr. Kamakshinatha Sivakumar, Technical Expert for the UNFPA PQ Programme</i>
11:00–11:30	Coffee/Tea break

11:30–12:00	New and innovative technologies for non-systematic contraceptives and STI barrier prophylactics . <i>Dr. Kamakshinatha Sivakumar, Technical Expert for the UNFPA PQ Programme</i>
12:00–12:45	Highlights of impact assessment of the WHO/UNFPA prequalification program <i>Speaker : Linda Serwaa, Head Product Quality Assurance , UNFPA</i>
12:45–13:00	Q&A
13:00–14:00	Lunch
SESSION 6 – PARALLEL TRACK 14:00 – 17:00	WHO INSPECTION SERVICES <i>Session Chair: Stephanie Croft. Technical Officer, PQT/INS, WHO</i> <i>Attendance mode: Hybrid</i>
14:00–14:05	Introduction to PQT/INS programme of activities
14:05–14:30	PQT/INS Programme update (overall)
14:30–14:45	Trends in the inspection of Medicines manufacturing sites (APIs and FPPs)
14:45–15:00	Trends in the inspection of CRO(s)
15:00–15:15	Trends in the inspection of Vaccine manufacturing sites
15:15–15:30	Trends in the inspections of Diagnostics
15:30–16:00	Coffee/Tea break
16:00–16:15	Trends in the inspection of Vector Control manufacturing sites
16:15–16:30	New IT system and impact on communication/data management
16:30–17:15	Updates on recent guidance published and impact on inspections
SESSION 7 – PARALLEL TRACK 14:00- 17:00	LOCAL PRODUCTION & TECHNICAL ASSISTANCE <i>Session Chair: Jicui Dong, Unit Head, LPA, WHO</i> <i>Attendance mode: Hybrid</i>
14:00-14:15	Opening session Updates on WHO Local Production and Assistance activities <i>Speaker: Jicui Dong, Unit Head, LPA, WHO</i>
14:15-14:25	WHO PQ/EUL specialized technical assistance: overview <i>Speaker: David Woo, Technical Officer, LPA, WHO</i>
14:25-14:35	Specialized technical assistance for medicines <i>Speaker: Kim Notenboom, Technical Officer, LPA, WHO</i>
14:35-14:45	Specialized technical assistance for vaccines <i>Speaker: Alan Fauconnier, Technical Officer, LPA, WHO</i>
14:45-14:55	Specialized technical assistance for IVDs <i>Speaker: David Woo, Technical Officer, LPA, WHO</i>
14:55-15:00	Closing remark

	<i>Speaker: Jicui Dong, Unit Head, LPA, WHO</i>
15:30–16:00	Coffee/Tea break
18:00–20:00 UN CITY CAFETERIA	RECEPTION FOR ALL PARTICIPANTS

DAY 3	WEDNESDAY, 29 NOVEMBER
08:00–08:30	ENTRANCE TO UN CITY
SESSION 8 – PLENARY: 08:30–13:00	PROCUREMENT UPDATES <i>Session chair: Francisco Blanco, Chief, Medicines and Nutrition, UNICEF</i> <i>Attendance mode: Hybrid</i>
08:30–08:45	Introduction to Procurement Update Session <i>Francisco Blanco - Chief, Medicines and Nutrition Center (MNC,) UNICEF</i>
08:45–09:00	WHO Procurement <i>Shipra Sharma – Procurement Officer, Supply Catalogues & LTAs, WHO</i>
09:00-09:45	Global Fund Procurement & Supply Updates <i>Clarisse Morris - Specialist, Global Sourcing Health Technologies</i>
09:45-10:00	Promoting access to TB products; GDF’s end to end approach <i>Magaly Babayel</i>
10:00–10:15	UNFPA Procurement and Strategic Sourcing <i>Roberto Mena - Head of Strategic Sourcing Team</i>
10:15-10:45	Coffee/Tea break
10:45-11:00	UNICEF Procurement: Comprehensive supply for impactful and cost-efficient health programmes <i>Cynthia Kamtengeni - Contracts manager</i>
11:00-11:15	UNHCR Procurement <i>Federico Pasqualini</i>
11:15–11:30	UNDP Global Health Procurement and Supply chain architecture 2023 <i>Zafar Yuldashev</i>
11:30-11:45	PAHO Procurement <i>Jordi Balleste, Strategic Fund Procurement - Unit Chief, Procurement and Supply Management, PAHO</i>
11:45-12:00	Sustainable Procurement- high level messages to manufacturers and suppliers <i>UNICEF (CC/MSFIC/QAC – Silvia Uneddu)</i>
12:00-12:30	Plenary and Q&A
12:30-13:30	Lunch Break

SESSION 9 – PARALELL TRACK 13:30–17:00		DEEP DIVE INTO UNICEF SPECIFIC ISSUES <i>Session Chair: Cynthia Kamtengeni, Contracts manager, UNICEF</i> <i>Attendance mode: Hybrid</i>	
13:30-13:50	UNICEF programme and supply priorities 2023-2025 <i>Francisco Blanco, Chief, Medicines and Nutrition Center, UNICEF</i>		
13:50-14:30	Updates of UNICEF quality requirements and tools, Demonstration of Supplier Document Library <ul style="list-style-type: none"> • <i>Peter Ikamati, Technical Specialist</i> • <i>Rennie SC, Technical associate</i> • <i>Peter S Jakobsen, Quality Assurance Specialist</i> 		
14:30-15:00	Common quality issues- Case studies from dossier assessments, GMP inspections, complaints and recalls <ul style="list-style-type: none"> • <i>Rajiv Kshirsagar, Technical Specialist,</i> • <i>Rennie SC, Technical Associate</i> • <i>Unine Felix Quality Assurance Specialist</i> • <i>Kasra Ghasemi, Quality Assurance Specialist</i> 		
15:00-15:30	Coffee/Tea break		
15:30-16:00	Common issues in procurement and contract management- Case studies from LTAs, POs, GR, FF <i>Caroline Kiyiika/ Shahbaz</i>		
16.00-16.20	New Packing-Packaging-Labeling and barcoding Requirements for shippers and pallets <i>WIMC/ SCSC/MNC (Patrick Adler)</i>		
16:20-16:40	New Bid Declaration Form <i>UNICEF CC Policy Unit (Silvia Uneddu/Nor)</i>		
16:40-17:00	New Invoicing System <i>FMAC (Khounkham Seebounhouang)</i>		
17:00-17:30	Q&A		
SESSION 10: PARALELL TRACK 13:30–17:00		UNFPA's SUPPLY CHAIN MANAGEMENT: Common issues in procurement and contract management <i>Session Chair: Udara Bandara, Deputy Chief, SCMU, UNFPA</i> <i>Attendance mode: Hybrid</i>	
13:30-13:50	Common technical and solicitation issues related to Dossier submissions and dossier assessments. <ul style="list-style-type: none"> • <i>Cristina Palau; Contract Analyst, Supply Chain Management Unit, UNFPA.</i> • <i>Olga Maria Pineda Velasquez Technical Consultant UNFPA</i> 		
13:50-14:20	Contract management; case studies from LTAs, <i>Roberto Mena, Head, Strategic sourcing team</i>		

14:30-15:00	Management of Quality complaints, recalls and incidences; case studies <i>Linda Serwaa, Head, Product Quality Assurance</i>
15:00-15:30	Coffee/Tea break
15:30-16:00	Prepositioning and inventory management: UNFPA 's approach, common issues and challenges encountered <i>Daniela Andries, Logistics and Inventory Management Specialist Supply Chain Management Unit</i>
16.00-16.30	Emergency procurement with its associated challenges <i>Andres Blasco, Procurement Specialist, UNFPA Supply Chain Management Unit</i>
16:30-17:00	Q &A

DAY 4	THURSDAY, 30 NOVEMBER
07:00–08:40	ENTRANCE TO UN CITY
SESSION 11 – PLENARY SESSION: 08:45–13:00	REGULATORY UPDATES <i>Session Chair: Hiiti Sillo, Unit Head, Regulation and Safety, WHO</i> <i>Attendance mode: Hybrid</i>
8:45-10:30	<p>Introductions and objectives of the session <i>Hiiti Sillo, Unit Head, Regulation and Safety, WHO (5 minutes)</i></p> <p>The new era of WLA (15 minutes) <i>Alireza Khadem, Team Lead REG/RSS (virtually)</i></p> <p>Reliance and Facilitated Regulatory Pathways including CRP (15 minutes) <i>Marie Valentin, Team Lead, REG/FPI</i></p> <p>Pannel discussion 1h10 (presentation 10 minutes each and then panel discussion and questions from the audience)</p> <p>Industry point of view – best use of reliance and WLA and Regulators journey to WLA</p> <ul style="list-style-type: none"> • Ms. Prisha Patel Prisha Patel, Senior Manager, International Regulatory Science & Policy Global Product Development, Pfizer, UK on behalf of IFPMA, • Mr Parag Nagarkar, Head of International Regulatory Affairs, Serum Institute of India will join the panel on behalf of SIPL/DCVMN (virtually). • Dr Youngjin Ahn, Director of Pharmaceutical Policy Division, Ministry of Food and Drug Safety (virtually).
10:30-11:00	Coffee/Tea break
11:00-13:00	<p>Introductions and objectives of the session <i>Hiiti Sillo, Unit Head, Regulation and Safety, WHO (5 minutes)</i></p> <p>Safety Monitoring (15 minutes) <i>Speaker: Shanthi Pal, Team Lead, REG/PVG (virtually)</i></p>

	<p>Supply Chain Integrity – updates on DEG / EG contamination of syrup medicines) (15 minutes) <i>Speaker: Rutendo Kuwana, Team Lead, REG/ISF (virtually)</i></p> <p>Nitrosamine Exchange – A Global Knowledge base (20 minutes) <i>Speaker: Frederick Meadows, USP</i></p> <p>Norms and Standards for Product Regulation (15 minutes) <i>Speaker: Steve Esteveao Cordeiro, Technical Officer, HPS/NSP</i></p> <p>Updates on work from the Expert Committee on Biological Standardization <i>Speaker: Ivana Knezevic, Team Lead, HPS/NSB (15 minutes) (virtually)</i></p> <p>Questions and answers (35 minutes)</p>
13:00–14:00	Lunch
SESSION 12 – PARALLEL SESSION: 14:00–16:00	TREATMENT AND DIAGNOSTIC GUIDELINES, ESSENTIAL MEDICINES LISTS (Inc for Children), ESSENTIAL IN VITRO DIAGNOSTICS LIST <i>Moderator: Steve Estevão Cordeiro, Technical Officer, WHO/NSP</i> <i>Attendance mode: Hybrid</i>
14:00	Introductory remarks: <i>Clive Ondari, Director, WHO Department of Health Product Policy and Standards,</i>
14:05-14:15	Tuberculosis Guidelines <i>Matteo Zignol, Unit Head, WHO TB Prevention, Diagnosis, Treatment, Care & Innovation Unit</i>
14:15-14:25	HIV and STIs <i>Marco Vitoria, Medical Officer, Treatment and Care Team, WHO Global HIV, Hepatitis and Sexually Transmitted Infections Programmes</i>
14:25-14:35	COVID therapeutic guidance and process for new influenza guidelines <i>Janet Diaz, Network Leader, WHO Health Care Readiness</i>
14:35-14:45	Malaria <i>Silvia Schwarte; Technical Officer; Diagnostics, Medicines and Resistance Unit; WHO Global Malaria Programme</i>
14:45-14:55	Hepatitis <i>Philippa Easterbrook, Scientist, WHO Global HIV, Hepatitis and Sexually Transmitted Infections Programmes</i>
14:55-15:05	WHO Model List of Essential Medicines <i>Lorenzo Moja, Technical Officer, WHO EML Secretariat</i>
15:05-15:15	Fourth WHO model list of essential in vitro diagnostics <i>Ana Aceves, Technical Officer, WHO EDL Secretariat</i>
15:15-15:30	Q&A
SESSION 13: PARALLEL SESSION 14:00 – 16:00	REQUIREMENTS FOR STORAGE AND TRANSPORTATION OF TIME AND TEMPERATURE SENSITIVE HEALTH PRODUCTS (TEMPERATURE CONTROL AND MONITORING OF SHIPMENTS) <i>Chair: Dimitrios Catsoulacos, Technical Officer, PQT/INS, WHO</i>

<i>Attendance mode: Hybrid</i>	
14:00 – 16:00	<ul style="list-style-type: none"> • How the UN 3PL intermodal freight forwarder strategy managed by UNICEF on behalf of the UN is aligned to a decentralization strategy – <i>Jean Cedric Meeus, Chief, Global Transport Center, UNICEF SD</i> • UNFPA presentation of the temperature monitoring program – <i>Johnson Moyo, Quality Assurance Associate & Linda Serwaa, Head, Quality Assurance Associate, UNFPA Product Quality Assurance team.</i> • International Logistics Management- UNFPA's logistics operations – <i>Daniela Andries & Joanna Trachimowicz, UNFPA logistics unit.</i> <p style="text-align: right;">Q & A</p>
SESSION 14 - PLENARY 16:00 – 17:00	PLENARY DISCUSSION & OFFICIAL MEETING CLOSURE <ul style="list-style-type: none"> • Key observations and thanks • Closing remarks by Dr Rogerio Gaspar, Director, RPQ, WHO

DAY 5	FRIDAY, 1 DECEMBER 2023
1-TO-1 MEETINGS – WITH PARTICIPATING AGENCIES	
<i>NOTE: 1-To-1 Meetings will not only be held on Day 5, but they can be scheduled on any day starting from Day 1. Participants are encouraged to contact the agency of interest below to be able to schedule a meeting between Day 1 and 5 as schedules allow</i>	
<i>Entrance to UN City for a 1-to-1 meeting will be dependent on meeting confirmation (including meeting time and location) from the agency with whom the meeting has been requested.</i>	
Meeting participants can request a meeting by contacting these agency staff: <ul style="list-style-type: none"> • For WHO Prequalification Inspection Services: Stephanie Croft, crofts@who.int • For WHO Prequalification of In Vitro Diagnostics: Irena Prat prati@who.int • For WHO Prequalification of Medicines: Matthias Stahl stahlm@who.int • For WHO Prequalification of Vaccines & Immunization Devices: Carmen Rodriguez Hernandez rodriguezhernandezc@who.int • For WHO Prequalification of Vector Control Products: Dominic Schuler, schulerd@who.int • For WHO Local Production: Jicui Dong dongji@who.int • For WHO Regulatory Updates: Marie Valentin, valentinm@who.int • For UNICEF Inspection Services; Helene Moller; hmoller@unicef.org • For UNICEF Medicines Francisco Blanco fblanco@unicef.org and Mary Atieno Ojoo Mary Atieno Ojoo@unicef.org • For UNICEF Vaccines; Ann Ottosen; aottosen@unicef.org, • For UNICEF In Vitro Diagnostics, Vector control products Wandani Sebonego wsebonego@unicef.org; • For UNICEF Medical devices and consumables; Selenge Lkhagva slkhagva@unicef.org • For UNICEF Cold Chain and Immunization devices; Thomas Sorensen tsorensen@unicef.org; • For UNFPA: Linda Serwaa serwaa@unfpa.org; • Ashley Moyo asmoyo@unfpa.org ; Mona Ghasemi ghasemi@unfpa.org 	