

Provisional agenda

- 1. Opening of the meeting**
- 2. Adoption of the agenda and programme of work**
- 3. Update by the Secretariat on the activities and budget to implement the workplan of the Member State mechanism**
- 4. Update on the list of prioritized activities for 2022–2023**
 - (A) Strengthen the capacity of national/regional regulatory authorities for the prevention and detection of, and response to, substandard and falsified medical products
 - (B) Develop, expand and maintain global networks of stakeholders to facilitate cooperation and collaboration
 - (C) Improve Member States' understanding and uptake of technologies to screen and detect substandard and falsified medical products, and the implementation of national traceability systems
 - (D) Leverage the competencies of relevant stakeholders, including policy-makers, procurers, distributors, practitioners, patients and consumers, and good governance to reduce the burden of substandard and falsified medical products
 - (E) Enhance Member States' capacity to run effective risk communication campaigns for substandard and falsified medical products
 - (F) Enhance Member States' capacity to expand awareness, effectiveness, impact and outreach in their work on substandard and falsified medical products
 - (G) Identify and develop appropriate strategies to understand and address the distribution or supply of substandard and falsified medical products via the internet
 - (H) Develop strategies for national regulatory authorities to mitigate public health risks posed by the distribution of substandard and falsified medical products through informal markets
- 5. WHO's participation in relevant global and regional initiatives**
- 6. Future work of the Member State mechanism**

- 7. Proposed dates of the twelfth meeting of the Member State mechanism**
- 8. Report of the Member State mechanism**
- 9. Closure of the meeting**

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