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