

Provisional agenda

- 1. Opening of the meeting**
- 2. Adoption of the agenda and method 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 2018–2019**
 - (A) Develop and promote training material and guidance documents to strengthen the capacity of national/regional regulatory authorities for the prevention and detection of and response to substandard and falsified medical products
 - (B) Expand and maintain the global focal point network among national medicines regulatory authorities to facilitate cooperation and collaboration
 - (C) Improve understanding of Member States on detection technologies, methodologies and “track and trace” models
 - (D) Increase Member States’ knowledge of the links between substandard and falsified medical products and access to quality, safe, efficacious and affordable medical products
 - (E) Develop and leverage existing activity for effective risk communication and make recommendations for awareness campaigns on 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) Promote shared understanding among Member States from a public health perspective regarding medical products in transit
 - (H) Identify and develop appropriate strategies to understand and address the distribution or supply of substandard and falsified medical products via the internet
- 5. WHO participation in the Global Steering Committee for Quality Assurance of Health Products**
- 6. Update on WHO activities for regulatory systems strengthening**
- 7. Update on governance issues**

- 8. Proposed dates of the eighth meeting of the Member State mechanism**
- 9. Report of the Member State mechanism**
- 10. Closure of the meeting**

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