01 June 2020

Member State Mechanism on Substandard and Falsified (SF) Medical Products

Meeting of the Steering Committee
24-25 June 2020
Virtual meeting

Draft Agenda

1. Opening of the meeting and adoption of the agenda

2. Update by the Secretariat on activities and budget to implement the work plan of the Member State Mechanism

3. Update of the list of prioritized activities for 2020-2021

    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 SF medical products

    B. Expand and maintain the Global Focal Point Network among national medicines regulatory authorities to facilitate cooperation and collaboration

    C. Improve Member States' understanding of detection technologies, methodologies, and "track and trace" models

    D. Increase Member States' knowledge of the links between SF 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 SF medical products

    F. Enhance Member States' capacity to expand awareness, effectiveness, impact and outreach in their work on SF 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 SF medical products via the internet

4. Update on WHO's participation in relevant global and regional initiatives

5. Planning and logistics for MSM meetings in 2020, including an update on governance issues

6. Closure