

Substandard and falsified medical products

The Seventy-sixth World Health Assembly, having considered the consolidated report by the Director-General,¹

Decided to request the Director-General:

- (1) to facilitate the conduct of an independent evaluation of the Member State mechanism on substandard and falsified medical products in accordance with the terms of reference to be developed by the Steering Committee of the Member State mechanism;
- (2) to report on the outcome of the evaluation to the governing bodies consistent with existing reporting requirements of the Member State mechanism on substandard and falsified medical products.

Ninth plenary meeting, 30 May 2023
A76/VR/9

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¹ Document A76/7 Rev.1.