HTA CG | MEMBER STATE COORDINATION GROUP ON HEALTH TECHNOLOGY ASSESSMENT

Short introduction to HTAR

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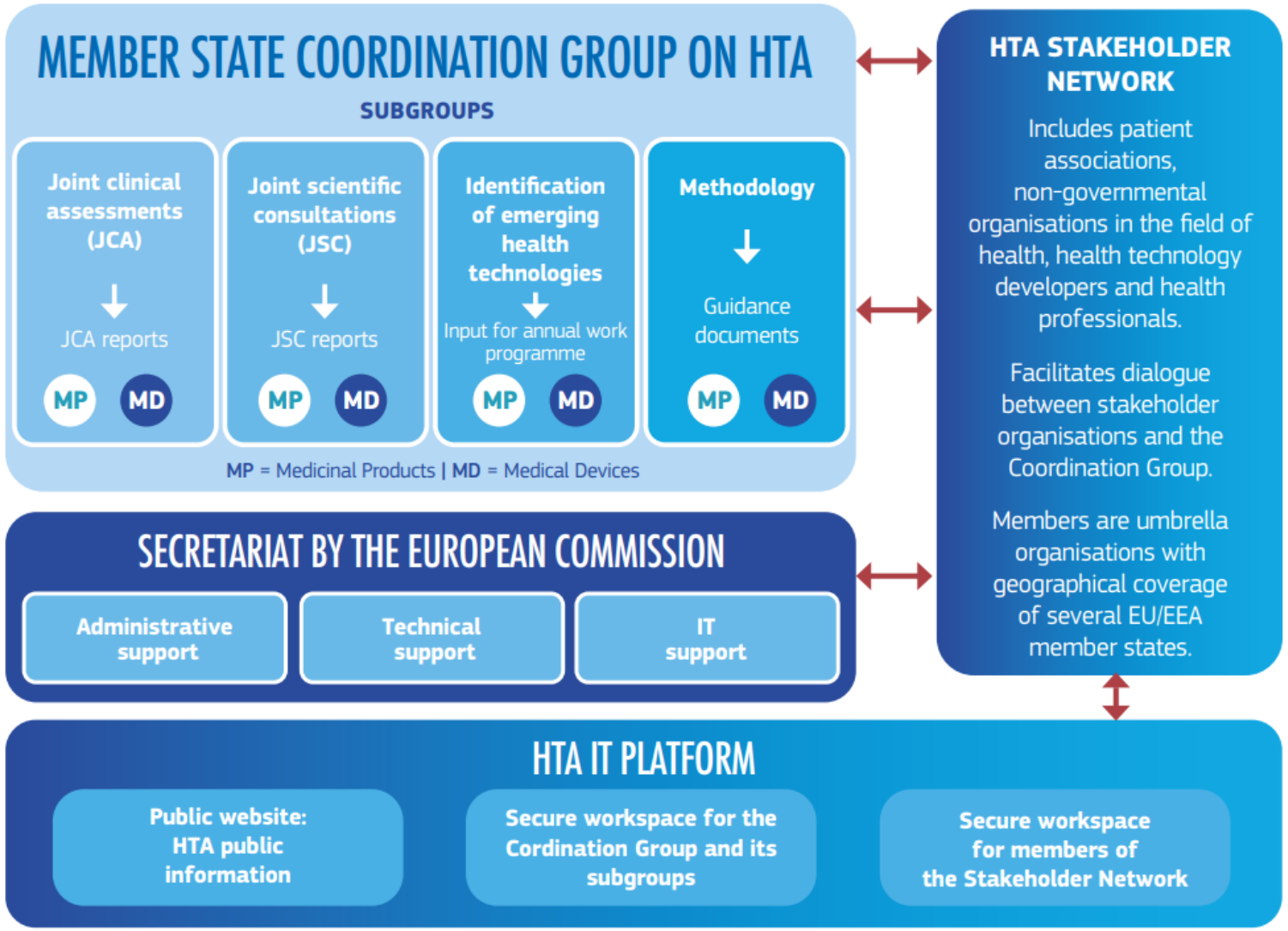
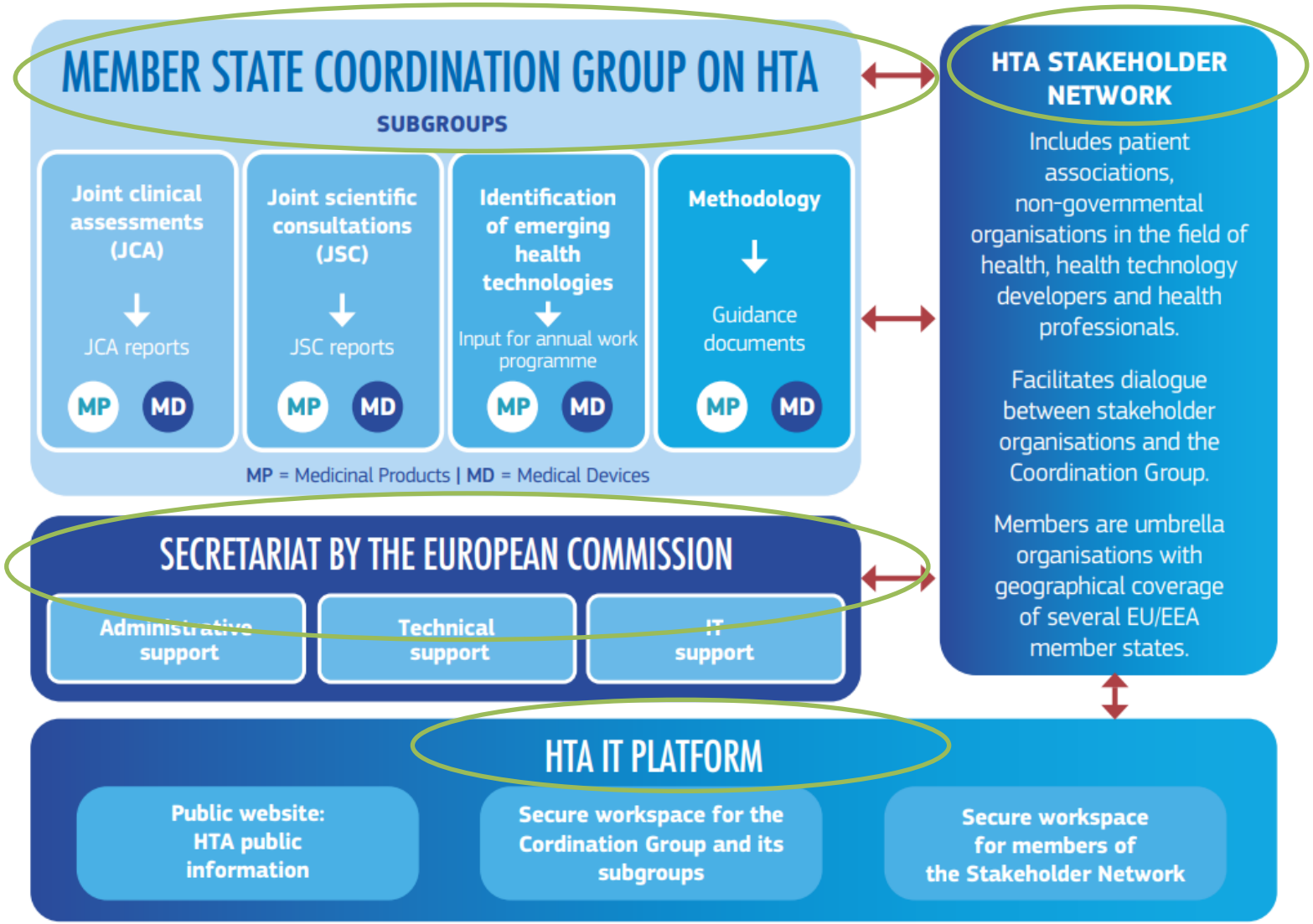
Governance


MEMBER STATE COORDINATION GROUP ON HTA
SUBGROUPS
Joint clinical assessments (JCA)
JCA reports
MP
MD
Joint scientific consultations (JSC)
JSC reports
MP
MD
Identification of emerging health technologies
Input for annual work programme
MP
MD
Methodology
Guidance documents
MP
MD
MP = Medicinal Products | MD = Medical Devices
HTA STAKEHOLDER NETWORK
Includes patient associations, non-governmental organisations in the field of health, health technology developers and health professionals.
Facilitates dialogue between stakeholder organisations and the Coordination Group.
Members are umbrella organisations with geographical coverage of several EU/EEA member states.
SECRETARIAT BY THE EUROPEAN COMMISSION
Administrative support
Technical support
IT support
HTA IT PLATFORM
Public website: HTA public information
Secure workspace for the Cordination Group and its subgroups
Secure workspace for members of the Stakeholder Network

Joint Configuration

First Point of Contact

Public Platform Europa website



Meet biannually F2F

HTAR – 3 years on

- 21 Guidance, Procedures and templates published (HTACG)
- 6 Implementing Acts (Member States)
- IT Platform developed to conduct joint work (EC)

These are the resources you will need when submitting under the HTAR

Information on work of HTACG and Subgroups

Member State Coordination Group on HTA (HTACG)

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The Regulation on health technology assessment (HTAR) established the Coordination Group on Health Technology Assessment (the 'HTACG') composed of Member States' representatives, mainly from HTA authorities and bodies.

The key tasks of the HTACG are to coordinate and adopt the joint HTA work carried out by its sub-groups within the scope of this Regulation and to adopt methodological and procedural guidance documents for joint work.

The HTACG also aims to ensure cooperation between the relevant European Union bodies (e.g. the European Medicines Agency), as well as appropriate involvement of stakeholder organisations and experts in its work.

Members of the HTACG

The members of HTACG are designated by the Member States in accordance to Article 3(2) of the Regulation (EU) 2021/2282, following a request from the Commission. Observers from EEA countries are designated following a similar procedure.

As laid down in Article 3(8), the HTACG will provide expertise on HTA for both medicinal products and medical devices (the latter covering also in vitro diagnostic medical devices).

- Members and EEA observers of the HTACG – medical devices
- Members and EEA observers of the HTACG – medicinal products

Members of the subgroups

https://health.ec.europa.eu/health-technology-assessment/implementation-regulation-health-technology-assessment/member-state-coordination-group-hta-htacg_en