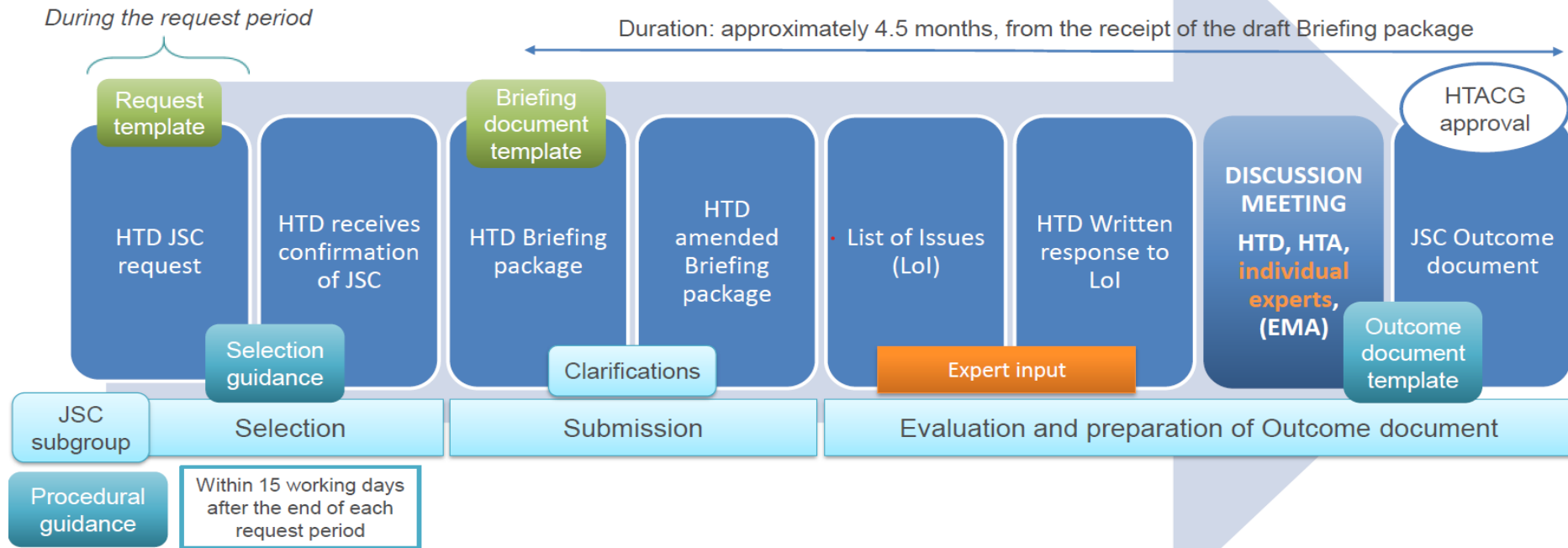


JSC process overview



Source: HTACG presentation at the webinar for HTDs 21.03.2025

Medicinal Products (MP)	Medical Devices (MD) and In Vitro Diagnostic Medical Devices (IVD)
Procedural Guidance for JSC on MP – adopted on 28.11.2024	Procedural Guidance for JSC on MD and IVD – adopted on 15.04.2025
Submission request template for HTACG JSC and Parallel HTACG/EMA JSC for MP – adopted on 28.11.2024	Submission request template for the HTACG and Parallel HTACG/Expert Panels JSC – adopted on 15.04.2025
Guidance for the selection of MP for JSC – adopted on 28.11.2024	Guidance for the selection of MD and IVD for JSC – adopted on 15.04.2025
Briefing document template for JSC for MP – adopted on 28.11.2024	Briefing Document template for MD – adopted on 28.02.2025
Briefing document template for Parallel HTA HTACG/EMA JSC for MP – adopted on 28.11.2024	Briefing Document template for IVD – adopted on 28.02.2025
	Briefing Document template for parallel HTACG/Expert Panels Joint Scientific Consultations for medical devices – adopted on 09.06.2025
Outcome document for JSC on MP – adopted on 28.11.2024	Outcome document template for MD and IVD – adopted on 28.02.2025 r.

Abbreviations	Definitions
JSC	Joint Scientific Consultations
HTD	Health Technology Developer
HTACG	The Health Technology Assessment Coordination Group established under Regulation (EU) 2021/2282
EMA	European Medicines Agency