HTA Expert meeting

Current experience and developments in HTA of medtech[1] in Europe

8 May 2014 between 10:00 and 17:00 at Eucomed / EDMA offices (Rue Joseph II, 40 – 1000 Brussels)

9:30 – 10:00	**Registration and Coffee**
10:00 – 10:15	**Welcome and Introduction**
10:15 – 11:30	**HTA for medtech: For different medical technologies, where does HTA fit in the care pathway in Europe? (INDUSTRY PRESENTATIONs)** • Pharma vs. medical technology models • Examples of the role of HTA in the healthcare pathway for medtech **Presentation to cover examples such as:** • Reimbursement pathways and reimbursement consequences • Different methods and sources of information • Innovation cycles and selection of technologies for HTA • HTA before or after launch, periodicity of HTA • Data requirements to demonstrate value
11:30 – 12:30	**WP7 SG3 Methodological Guideline on Medical Devices. The current guideline development process in EUnetHTA and guideline structure (EUnetHTA PRESENTATION)** **Discussion with industry** • Input in the form of general comments and recommendations for special contents and methodological aspects
12:30 – 13:15	**LUNCH**
13:15-15:00	**EUnetHTA WP4 and WP5 Strand B: Discussion on challenges and current processes (EUnetHTA PRESENTATION)** • Application/transferability of deliverables to national/local level HTA • Identification of all manufacturers, identification and selection of other stakeholders, best ways to contact the industry • Tasks of the industry (evidence submission, other relevant information) • Scoping meeting(s), WP4 suggestion • Confidentiality issues (evidence, confidential version of assessment)

[1] medtech = Medical Devices, In-vitro Diagnostics and Medical Imaging

	WP4 and WP5 Strand B experience (INDUSTRY PRESENTATIONS) • WP4: EDMA experience • WP5 Strand B: Eucomed experience (Renal Denervation example) **Presentations to cover:** • Limitations, including topic selection process and criteria • How to address limitations • Potential way forward
15:00 – 15:15	**Coffee break**
15:15 – 16:15	**WP7 SG4: Template development for medical devices** (excluding IVDs) **(EUnetHTA PRESENTATION)** Discussion about the analysis and draft template: WP7 SG4 will send a draft report of the findings of national evidence requirements for devices and an outline for a submission template. • Evidence requirements received from reimbursement agencies • Approach to developing submission template for devices • Introduction to draft template **Discussion with industry:** • Are our analyses complete? • Is there anything we are missing? • Are there any issues with any of the evidence requirements?
16:15 – 16:45	**WP7 SG1 Early Dialogues: Initial exchange of views and identification of the issues (EUnetHTA PRESENTATION)** **Discussion with industry:** • Concept • Expected benefits and possible barriers and limitations • Stakeholders to be involved • Way forward
16:45 – 17:00	**Wrap up, Conclusions and Next Steps**