

Joint Action on Market Surveillance of medical devices (JAMS)

The European Commission has made funding available to a group of European countries that agreed to participate in a three-year project which commenced in October 2016. This project aims to find ways for regulators of medical devices in Europe to improve the checking and monitoring of medical devices.

Aims

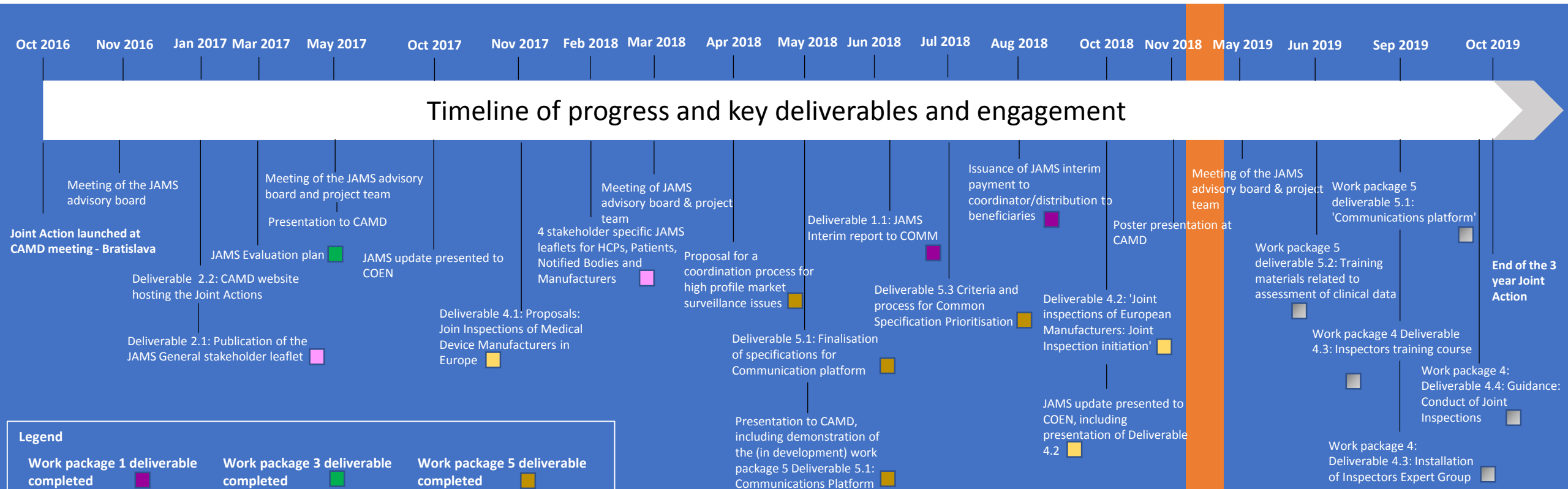
- To reinforce market surveillance between member states across Europe.
- Share and develop best practice, training, knowledge, and resources to increase public health protection in the medical devices sector.
- Improve coordination and help to develop skills and capacity in the market surveillance network.
- Ensure a consistent and proportionate approach to the work of manufacturer inspections.
- Identify and develop training needs for the clinical assessment of medical devices.

Participants

UK (MHRA)	CYPRUS (CYMDA)
IRELAND (HPRA)	CZECH REPUBLIC (MZCR)
FRANCE (ANSM)	DENMARK (DKMA)
NETHERLANDS (IGZ)	ESTONIA (TERVISEAMET)
AUSTRIA (AGES)	GERMANY (BFARM)
BELGIUM (FAMHP)	ITALY (SANITA)
CROATIA (HALMED)	LATVIA (VI)
	NORWAY (HESEDIR)
	PORTUGAL (INFARMED)
	SPAIN (AEMPS)
	SWEDEN (MPA)

Structure

- Work package 1: Coordination of the project (Led by UK)
- Work package 2: Dissemination of information (Led by UK)
- Work package 3: Evaluation of the project (Led by UK)
- Work package 4: Joint inspections of manufacturers (Led by Ireland & France)
- Work package 5: Clinical process and resources development (Led by Ireland)



Legend

Work package 1 deliverable completed	Work package 3 deliverable completed	Work package 5 deliverable completed
Work package 2 deliverable completed	Work package 4 deliverable completed	Expected completion of deliverable

Website: <https://www.camd-europe.eu/joint-action-projects/market-surveillance-of-medical-devices-jams/>

