Valvira

	RoHS Directive 2011/65/EU, medical devices
	National Supervisory Authority for Welfare and Health (Valvira)
Surveillance	1. Reactive market surveillance 2. Proactive market surveillance - Finnish medical device manufacturers (document inspections) - Operators (document inspections) 3. Projects - Individual product groups
Communications	1. Keeping the Valvira website up-to-date 2. Providing information about RoHS requirements
Development	1. Cooperation with stakeholders (Ministry of the Environment, Tukes) 2. Joint projects of EU surveillance authorities if relating to MDs
Training	1. Manufacturer and operator training in the requirements set by the Directive (MD)
Other	If necessary, cooperation with electrical and environmental safety actors.