Commission Implementing Regulation (EU) No 920/2013 of 24 September 2013 on the designation and the supervision of notified bodies under Council Directive 90/385/EEC on active implantable medical devices and Council Directive 93/42/EEC on medical devices (Text with EEA relevance)

Article 1	Definitions
Article 2	Interpretation of designation criteria
Article 3	Procedure for the designation of notified bodies
Article 4	Extension and renewal of designation
Article 5	Surveillance and monitoring
Article 6	Investigation of the competence of a notified body
Article 7	Exchange of experience on designation and supervision of conformity assessment bodies
Article 8	Operating of designating authorities
Article 9	Cooperation with accreditation bodies
Article 10	Entry into force and date of application
	Signature

ANNEX I

Interpretation of the criteria set out in Annex 8 to Directive 90/385/EEC and in Annex XI to Directive 93/42/EEC

1. Sections 1 and 5 of Annex 8 to Directive 90/385/EEC...

2. The second paragraph of Section 2 of Annex XI to...

3. Sections 3 and 4 of Annex 8 to Directive 90/385/EEC...

4. Sections 6 of Annex 8 to Directive 90/385/EEC and of...

5. Sections 7 of Annex 8 to Directive 90/385/EEC and of...

ANNEX II

Application form to be submitted when applying for designation as notified body

Designating authority: …
Name of the applying conformity assessment body: …
Previous name (if applicable): …
EU Notified Body number (if applicable): …
Address: …
…
…
…
Contact person: …
E-mail: …

Telephone: …
Legal form of the conformity assessment body: …
Company registration number: …
At company register: …
…
…
The following documents shall be added. In case of extension...

(**1**) OJ L 189, 20.7.1990, p. 17.

(**2**) OJ L 169, 12.7.1993, p. 1.

(**3**) OJ L 218, 13.8.2008, p. 30.