



Product Service

Appendix A ?

Application for performance of a conformity assessment procedure in accordance with 90/385/EEC (AIMD)



Manufacturer:



Application identification:

Appendix A

Device details

- AIMD 0101 Active implantable medical devices for stimulation / inhibition
- AIMD 0102 Active implantable medical devices delivering drugs or other substances
- AIMD 0103 Active implantable medical devices substituting or replacing organ functions

	? Name of model Trade name	? Model (number(s)) ^{*1)}	Name (from UMDNS)	? UMDNS Code ^{*2)}	? OEM Device	manufactured utilizing animal tissues or derivatives rendered non-viable	Incorporates a substance which if used separately may be considered to be a medicinal product as defined in Directive 2001/83/EC or a human blood derivative
1.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
2.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
3.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
4.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
5.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
6.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
7.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
8.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
9.					<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

^{*1)} This information is only required for product-related procedures (i.e. only for procedures in accordance with Annex 2.4, 3, 4).

^{*2)} The [UMDNS Code](#) is only required for product-related procedures for reporting to DIMDI.