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Zentralstelle der Länder  
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bei Arzneimitteln und  
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ZLG-BS-244.10.08



Product Service

# EC Certificate

Full Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)  
(Devices in Class IIa, IIb or III)

**No. G1 067972 0007 Rev. 01**

### Facility(ies):

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