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Zentralstelle der Länder
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bei Arzneimitteln und
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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)

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Facility(ies):

**Sichuan Nigale Biotechnology Co., Ltd.
No.28 Kuixing Road, 641400 Jianyang, Sichuan,
PEOPLE'S REPUBLIC OF CHINA**

**Sichuan Nigale Biotechnology Co., Ltd.
4th F, No.2 Factory Building, Shiyang Industrial
Park, No.55, Section 5th, Qingyun Village, Hi-Tech
District, 610041 Chengdu, PEOPLE'S REPUBLIC OF
CHINA**