EC Certificate Full Quality Assurance System: Certificate KR07/00886

The management system of

AMPall Co., Ltd.

3F, Annex HanKook Junja HyeopDong B/D
371-51, Gasan-Dong, Geumcheon-Gu, Seoul, Korea

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Infusion pump (Model: IP-7700);
Syringe pump (Model: SP-8800);
Blood Pressure Monitor (Model: BP 868F);
PCA pump (Model: PP-9900) and infusion set (PP-9900ACB series);
Sterile single use PCA pump (Model: PP-9800B1, PP-9800B2, PP-9800C1, PP-9800C2)

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 9 September 2013 until 12 July 2018 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 4 June 2016
Issue 8. Certified since 12 July 2007

Certification is based on reports numbered KR/SEL Y-PC/07167

Authorised by

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