

The management system of

Renacon Pharma (Private) Limited

18 - K.m. Ferozepur Road,
Lahore, Pakistan

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

Haemodialysis Concentrates.

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 22 January 2011 until 15 October 2013 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 15 October 2013
Issue 6. Certified since 15 October 2007

Certification is based on reports numbered GB/PI 216279

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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