EC Certificate Full Quality Assurance System: Certificate PK07/73231

The management system of

Renacon Pharma (Private) Limited

18 - K.m. Ferozepur Road, Opp Nishtar Colony
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
RENACARB (Part A: Solution; Part B: Powder);
RENACIT (Citrate Concentrate: Powder);
RENACIT –S (Citrate Concentrate: Part A: Solution; Part B: Powder)
RENACART (Bicarbonate Cartridge);
RENABAG (Bicarbonate Bag);
RENAPULV (Powder Concentrate);
RENAPULV-N (Powder Concentrate)
RENACATE (Acetate Concentrate Solution);
RENALACT (Lactate Concentrate
RENALACT (Lactate Concentrate)

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 10 October 2017 until 15 October 2021
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 15 October 2019
Issue 11. Certified since 15 October 2007

Certification is based on reports numbered PK/LHR 216279

Authorised by

SGS United Kingdom Ltd, Notified Body 0120
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com