SGS

EC Certificate Full Quality Assurance System: Certificate US15/842160

The management system of

Poriferous LLC

535 Pine Road,

Newnan GA 30263. United States 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

Sterile craniomaxillofacial polyethylene implants for reconstructive plastic surgery.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 13 May 2015 until 13 May 2020 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 19 March 2018

Issue 1. Certified since 13 May 2015

Certification is based on reports numbered WW/ME 606020

Authorised by



SGS United Kingdom Ltd, Notified Body 0120

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