EC Certificate Production Quality Assurance System: Certificate GB20/965119



The management system of

GSK Consumer Healthcare (GMDT) also trading as SmithKline Beecham Limited Slough

Clocherane, Youghal Road, Dungarvan, Co. Waterford, Ireland

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile single-use biopsy punches and curettes

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination

Certificate according to Annex III is a mandatory requirement for each device
in addition to this certificate to place that device on the market.

This certificate is valid from 15 January 2020 until 17 October 2023 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 17 October 2006 and first certified by SGS Belgium NV since 03 April 2019

Certification is based on reports numbered GB/PC/ 214615

Authorised by

SGS Belgium NV, Notified Body 1639

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