

The management system of

Super Seton B.V.

Paasheuvelweg 25
1105 BP Amsterdam, The Netherlands

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC on medical devices, Annex V

For the following products

SuperSeton: sterile seton drain for the treatment of perianal fistula.

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 18 November 2020 until 23 July 2023
and remains valid subject to satisfactory surveillance audits.
Issue 3. Certified since 19 December 2018.

Certification is based on reports numbered BE/AMD 17/1104.QMD

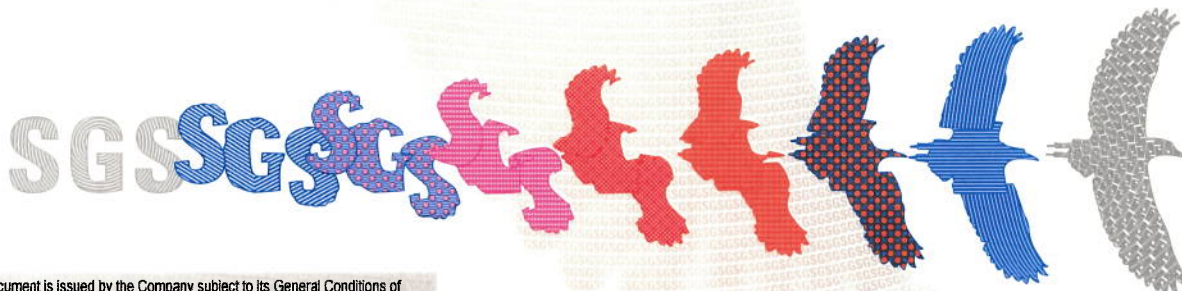
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