

METROVISION

4 rue des platanes
59840 Perenchie, France

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

**Vision Monitor:
device for testing visual functions based on electrophysiological,
psychophysical and video-oculography techniques.**

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 1 July 2015 until 18 March 2020 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 14 November 2017.
Issue 1. Certified since 1 July 2015.

Certification is based on reports numbered BE/AND 14/1211.QMD.

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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Page 1 of 1

