Adaptive Sensory Technology

DECLARATION OF CONFORMITY

Adaptive Sensory Technology GmbH
Maria-Goeppert-Str. 1
23562 Lübeck

hereby declares under their sole responsibility that the medical device

Manifold Contrast Vision Meter

meets the provisions of European Directive 93/42/EEC on medical devices.
This declaration is valid for three years and for the following devices:

Reference number	001.000.003 (comprising Server 001.001.003 and Remote Control 001.002.002/003)
Software version	2.6

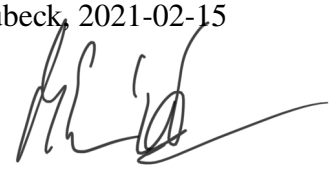
Any modification of the medical device not authorized by AST will invalidate this declaration.

The device was classified according to MDD Annex IX as active, not invasive, reusable, non-sterile and for short term use into

Class I following **rule 12.**

The conformity assessment procedure was performed according to MDD Annex VII.
A technical file detailing this declaration is kept.

Lübeck, 2021-02-15



Manuel Wille
(Managing Director of Adaptive Sensory Technology GmbH)