

EC Declaration of Conformity

To whom it may concern,

Samsø, Denmark, May 2024

We,

ExamVision ApS, Industrivej 11, Tranebjerg, 8305 Samsø, Denmark

hereby declare in our capacity as the product manufacturer, that the ExamVision Eyewear Panto which is intended to be used

as a visual aid for the Healthcare Professional while performing medical procedures,

complies with the requirements of the

REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No. 178/2002 and Regulation (EC) No. 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC with amendments.

Actor ID/SRN: DK-MF-000032397 **Basic UDI-DI:** 574400023EVEyewearDV

Product Name/Trade name: Panto

Intended Purpose: The ExamVision Eyewear is intended to be used as a visual aid

for the Healthcare Professional while performing medical

procedures.

Risk Class (MDR): Class I, rule I.

Conformity Assessment Route: Annex I General Safety and Performance Requirements

Annex II Technical Documentation

Annex III Technical Documentation on Post-Market Surveillance

Other relevant legislation, applicable

harmonised standards, and normative

documents:

RoHS ISO 12870:2018

ISO 21987:2017 ISO 14971:2019

Issue date: 1 May 2024

Signature:

Ole Anker Aagaard/Head of Legal