MONOQOOL

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EU DECLARATION OF CONFORMITY

Manufacturer: **MONOQOOL, Danstruplund, Danstrupvej 27M, DK – 3480 Fredensborg, SRN DK-MF-000025532**, in its capacity as the manufacturer hereby declares, under its sole responsibility, that the product listed below conforms to provisions of the Regulation (EU) 2017/745

Product: **Monoqool glasses frames** (in all their colors and sizes)
Classification: medical device class I

The product was classified as a class I medical device in conformity with Rule 1 Chapter III of Annex VIII of Regulation (EU) 2017/745.

We also declare that the product has been designed and manufactured under a quality management system compliant with Regulation (EU) 2017/745.

The product is in conformity with the requirements of Regulation (EU) 2017/745 and the standard EN ISO 12870:2016.

Product Details
Intended Purpose: The product is intended to hold prescription lenses used for correcting or improving the vision of patients with prescription glasses
Product Series: • Slider series • IQ mini series • IQ series • Wire series

Monoqool,
Allan G. Petersen, Founder

Responsible

Fredensborg, July 12, 2022

Place and Date