

EU Declaration of Conformity

Manufacturer:	Granberg AS Bjoavegen 1442, NO-5584 Bjoa, Norway	E-mail: post@granberg.no Phone: +47 53 775 300
Single Registration number (SRN):	NO-MF-00000207	Doc Nr: F-MD-003-018-1

REF. 210.0022E

The item is in conformity with (EU) 2017/745 Medical Device Regulation (MDR) as:

Name	Disposable Surgical Face Mask Granberg®
Description	Type IIR, non-sterile, blue
Risk Classification	Class I according to Rule 1 of Annex VIII of Regulation MDR (EU) 2017/745
Applied normative standards	EN ISO 13485:2016, EN ISO 10993-1:2020, EN ISO 10993-5:2009, EN ISO 10993-10:2023, EN ISO 10993-23:2021, EN ISO 14971:2019, ISO 15223-1:2021, EN ISO 20417:2021, EN 14683:2025
Intended Use	Medical face masks are designed to prevent transmission of infected agents, blood and/or bodily fluids from the person wearing them (including healthcare personnel and patients) to others.
Basic UDI-DI	702377GR210PPNSSM
Available sizes	One size fits all
EMDN Code	T020604

The technical documentation for assessing the conformity of the medical devices with MDR has been developed in accordance with Annexes II and III of MDR.

This Declaration of Conformity is issued under the sole responsibility of the manufacturer – Granberg AS.

Signed for Granberg AS:



Ole Marthon Granberg
Managing Director

Place and date of issue:

Bjoa, 10.11.2025