



## DECLARATION OF CONFORMITY

We, undersigned GIMA S.p.A., with operational headquarters in Gessate (MI), Via Marconi 1, and registered office in Milano, Via Tommaso Grossi 2, acting as manufacturer of the medical device:

GIMA Single Registration Number (SRN):

<b>Medical Device ( Trade Name and description)</b>	<b>Code</b>	<b>Basic UDI-DI code</b>
LUMINA LED LIGHT - trolley	49035	8023279Z120107020000000RC

Risk class I (Not sterile), according to the Rule 13 Annex VIII of Regulation (EU) 2017/745 (MDR), declares, under its own responsibility, that this medical device:

- comply with essential requirements and dispositions of Regulation (EU) 2017/745 (MDR), as from the Technical File filed at the company;
- common Specifications have not been used for the compliance of the above medical device;
- comply with directive 2011/65/EU (and subsequent amendments and integrations) on the restriction of the use of certain hazardous substances in electrical and electronic equipment.

Gessate, 5/28/2021

**GIMA S.p.A.**

The legal Representative  
(Nicola Manzoni)

A handwritten signature in black ink, appearing to read "N. Manzoni", is written over a horizontal line.