"EU" DECLARATION OF CONFORMITY

according to Regulation (EU) 2017/745, Annex IV

This declaration of conformity is issued under the sole responsibility of the manufacturer:

Product Source Jsc
51 Nguyen Trai, Ha Dong
Ha Noi, Viet, Vietnam

for the devices put into commerce under its own brand
DECLARES
that the device which this declaration is addressed to is conform to the
General Requirements for Safety and Performance required by annex I of Regulation (EU) 2017/745.

DEVICE NAME:	**Saline Solution**
MODEL:	**500ML**
TYPE:	**Medical Device**
CLASSIFICATION: (Rule 1. Annex Vill Regulation (EU) 2017/745)	**Class I**
BASIC UDI-DI:	**GTIN-8936226071819**
YEAR OF MANUFACTURE:	**2024**

Any modification applied to the medical device, if not expressively authorized by Empiry, cancel the validity of this declaration.

It's here declared that the device to which this declaration is addressed to, are conform to the following technical standards:

- **EN ISO 14971:2019+A11:2021**
- **EN ISO 10993-1:2010**
- **EN ISO 10993-5:2009**
- **EN ISO 10993-10:2013**

It's here declared that the device to which this declaration is addressed to, is commercialized in sterile package.

Signed for and on behalf of:

Registred Trade Mark: Nanocare

Strongbody UK Ltd.
40 Bowling Green Lane, London, EC1R ONE

Lion Rose - Director
Legal Representative

United Kingdom
25 April 2024

Legal Representative Signature