

## DECLARATION OF CONFORMITY

**Manufacturer: Informer Med sp. z o.o.**  
**Winogrody 118, 61-626 Poznań (Poland)**

Registration number SRN: PL-MF000003613

- **PRODUCT: Sterilization Pouch**
- **Classification: Class I non-sterile (accordingly to Annex VIII, chapter I MDR)**
- **Basic UDI-DI:** 5904305484020 – SSP0610, STERILIZATION POUCH 60mmx100mm  
5904305484211 – SSP0913, STERILIZATION POUCH 90mmx135mm  
5904305484228 – SSP0923, STERILIZATION POUCH 90mmx230mm  
5904305484235 – SSP1428, STERILIZATION POUCH 140mmx280mm  
5904305484242 – SSP2033, STERILIZATION POUCH 200mmx330mm  
5904305484259 – SSP2540, STERILIZATION POUCH 250mmx400mm  
5904305484716 – SSP3040, STERILIZATION POUCH 300mmx450mm  
5904305484723 – SSP1325, STERILIZATION POUCH 135mmx250mm  
5904305484730 – SSP3039, STERILIZATION POUCH 300mmx390mm
- **Certificate ISO 13485: in progress**

We declare that that the above mentioned products comply with The Regulation of the European Parliament and of the Council (EU) 2017/745 on medical devices (MDR), amendments to Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repeal of Council Directives 90/385/EEC and 93/42/EEC.

Date of issue: 25.05.2021

Signature:

*[Signature]*  
Inform Med Sp. z o.o.  
Członek Zarządu

*[Signature]*  
INFORMER MED Sp. z o.o.

*[Signature]*  
Arkadiusz Łeszyk  
Załącznik