

## EU DECLARATION OF CONFORMITY (No. 2019/252-01)

*This Declaration of Conformity is issued under the sole responsibility of the manufacturer*

### MANUFACTURER

<b>Company name</b>	Sterdoc B.V
<b>Full address</b>	Keersluisweg 21
<b>Postal code</b>	1332 EE
<b>Place</b>	Almere
<b>Country</b>	The Netherlands

### IDENTIFICATION MEDICAL DEVICE

<b>(Commerical) name</b>	Sterdoc
<b>Function/intended use</b>	Sterilization Packaging
<b>Type / model</b>	Reels, Tyveks and wrapping
<b>Batch/serial</b>	BT.FL/TY/WR.xxxx.050-1200
<b>Risk Class</b>	Class I (Rule I)

*The object of the Declaration described above is in conformity with all relevant provisions of*

**Medical Devices Directive 93/42/EEC**

*In conjunction with the following relevant harmonised standards or technical specifications*

**ISO-11607-1; ISO-11607-2**

*Signed for and on behalf of*

**Place of issue:** Almere  
The Netherlands

**Identity:** Peter Van der zee  
**Function:** Quality Manager

**Date of issue:** 1 December 2022

**Signature**

