



## Free Sales Certificate

The Danish Medicines Agency hereby certifies that the medical devices specified in the attached list are manufactured by:

**BiopSafe ApS**  
**Carolinevej 2A 1.tv**  
**DK-2900 Hellerup**

Medical devices which are CE marked in conformity with Directive 98/79/EEC meet the essential requirements for safety and performance. They may therefore be manufactured and marketed in Denmark and exported without any approval from the Danish Medicines Agency.



Valid from: 27 January 2022  
Valid Until: 27 January 2024

*Freja Bauch*  
**Freja Bauch**



We have been informed that the products specified below, are manufactured at the site:"

SP Medical A/S  
Møllevej 1  
DK-4653 Karise

"On behalf of"

BiopSafe ApS  
Carolinevej 2A 1.tv  
DK-2900 Hellerup

BiopSafe®, container and lid.



LÆGEMIDDELSTYRELSEN  
DANISH MEDICINES AGENCY