



Free Sales Certificate

The Danish Medicines Agency hereby certifies that the company has its registered place of business in Denmark:

BiopSafe ApS
Bygstubben 4
2950 Vedbaek
Denmark

Medical devices specified in the attached list which are CE marked in conformity with the Regulation on In Vitro Diagnostic Medical Device (Council Regulation 2017/746) may be manufactured and marketed in Denmark/EU and exported without any approval from the Danish Medicines Agency.



Valid from: 31 October 2023
Valid Until: 31 October 2025

Sultan Nur Bayram



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

SP Medical A/S
Møllevvej 1
4653 Karise
Denmark

"On behalf of "

BiopSafe ApS
Bygstubben 4
2950 Vedbaek
Denmark

BiopSafe®, container and lid.

Produktnavn / Product Name	Produkt ID / Product ID
BiopSafe® Biopsy Container 20ml	3178-200-xx
BiopSafe® Biopsy Container 60ml	3178-600-xx



LÆGEMIDDELSTYRELSEN
DANISH MEDICINES AGENCY