

**NOTARIUS PUBLICUS I KUNGSBACKA**  
NOTARY PUBLIC KUNGSBACKA, SWEDEN

**CERTIFICATE**

I, the undersigned, Joakim Gillersten, Notary Public in the City of Kungsbacka Sweden, do hereby certify that the attached document, under my seal of office, is a certified true photocopy corresponding with the original document.

Kungsbacka, this the 3<sup>rd</sup> day of December, 2021

Ex officio:

  
Joakim Gillersten  
Exp. No; 1466-2021



The Ministry for Foreign Affairs in Stockholm,  
hereby certifies that

**Joakim Gillersten**

Notary Public of Kungsbacka

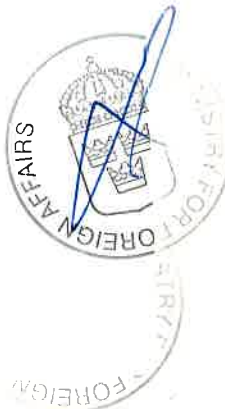
has issued and signed the foregoing  
attestation in his/her official capacity.

Stockholm 14 December 2021

Nr. 2982134

Fee Paid 180 SEK

  
Lisa Grehn



NOTARIUS PUBLICUS KUNGSBACKA

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CHỨNG NHẬN/HỢP PHÁP HÓA LÃNH SỰ  
CONSULAR AUTHENTICATION

1. Quốc gia ..... Việt Nam .....  
Country

Giấy tờ, tài liệu này  
This public document

2. do Ông (Bà) ..... Lisa Grehn ..... ký  
has been signed by

3. với chức danh  
acting in the capacity of

4. và con dấu của ..... Bộ Ngoại giao Trung Đức .....  
bears the seal/stamp of

được chứng nhận / hợp pháp hóa lãnh sự  
Certified

5. tại ..... Stockholm ..... 6. ngày 29.12.2021  
at

7. Cơ quan cấp ..... ĐSQ Việt Nam .....  
by

8. Số: 217/2021  
NP

TL. ĐẠI SỨ / FOR THE AMBASSADOR  
Bí thư thứ Hai / Second Secretary

  
Đào Tuấn Việt





# Quality System Certificate



Certificate No.:  
**DGM – 845**

Reference:  
**Aur1i2010v110f826**

Date of issue:  
**2021-02-02**

Valid Until:  
**2024-02-02**

Initial date of issue:  
**2014-09-30**

This is to certify that the quality system of:

**Vitrolife Sweden AB**  
**Gustaf Werners Gata 2**  
**SE-421 32 Västra Frölunda (visiting address)**  
**Box 9080, SE-400 92 Göteborg (postal address)**

fulfills the requirements in:

**EN ISO 13485:2016**

The certificate covers the following activities:

**Development, manufacture and marketing of products and systems for preparation, cultivation and storage of cells, intended for therapy**



The certificate is valid provided that the quality system continues to conform to the above-mentioned scope and provided that the company does not introduce substantial changes to the quality system without the approval of Presafe Denmark A/S. Certifications to DS/EN ISO 13485:2012, EN ISO 13485:2012, ISO 13485:2003, DS/EN ISO 13485:2016, EN ISO 13485:2016, ISO 13485:2016, DS/EN ISO 9001:2015, EN ISO 9001:2015, and ISO 9001:2015 include the requirements of any applicable corrigendum. The quality system certificate is issued pursuant to Presafe Denmark A/S' terms and conditions for the certification of quality systems for medical devices.

**Presafe Denmark A/S**  
Notified Body, Identification No. 0543  
Tuborg Parkvej 8, 2900 Hellerup, Denmark

  
Heidi Jørgensen

For Presafe Denmark A/S

