BUREAU VERITAS
Certification

VITREX Medical A/S

Vasekær 6-8, 2730 Herlev, Denmark

Bureau Veritas Certification Denmark A/S certifies that the Management System of the above organization has been audited and found to be in accordance with the requirements of the management system standards detailed below.

Standard

DS/EN ISO 13485:2016

Scope of certification

Development, manufacture and sales of single use glass and plastic capillary tubes and accessories for diagnostic use. Manufacture and sales of blood lancets and accessories for blood sampling and diagnostic. Marketing and sales of sutures, hypodermic syringes, Hypodermic needles, Cannulas, Needles and diabetic in-vitro medical devices.

Original cycle start date:	**14 June 2000**
Expiry date of previous cycle:	**NA**
Certification/Recertification Audit date:	**NA**
Certification/Recertification cycle start date:	**15 June 2018**

Subject to the continued satisfactory operation of the organization's Management System, this certificate expires on: **14 June 2021**

Certificate No.: DK009653 Version: 1 Revision date: **12 June 2018**

Certification Office: *Bureau Veritas Certification Denmark A/S*
Oldenborggade 25-31, 7000 Fredericia, Denmark

Further clarifications regarding the scope of this certificate and the applicability of the Management System requirements may be obtained by consulting the organization. To check this certificate validity, please call **(+45) 77 311 000.**

DANAK
SYSTEM Reg.nr. 5005

SERTIFIKATAS

VITREX Medical A/S

Vasekær 6-8, 2730 Herlev, Denmark

Bureau Veritas Cettification Denmark A/S ceñifies kad minėtos organizacijos valdymo sistema buvo patikrinta ir nustatyta, kad ji atitinka toliau nurodytus valdymo sistemos standartų reikalavimus.

Standartas

DS/EN ISO 13485:2016

Sertifikavimo apimtis

Vienkartinių stiklinių ir plastikinių kapiliarų vamzdžių ir jų priedų diagnostikos reikmėms kūrimas, gamyba ir pardavimas.

Kraujo lankstų ir aksesuarų, skirtų kraujo mėginių ėmimui ir diagnostikai, gamyba ir pardavimas. Siūlų, švirkštų, švirkštų, hipoderminių adatų,

Cannulas, adatos ir diabetiniai in-vitro medicinos prietaisai.

Originalo ciklo pradžios data: 2000 m. Birželio 14 d.

Ankstesnio ciklo galiojimo pabaiga:

Sertifikavimas / recertifikavimas Audito data:

Sertifikavimo / pakartotinio sertifikavimo ciklo pradžios data: 2018 m. Birželio 18 d

Atsižvelgiant į tolesnį patenkinamą organizacijos veiklą

Vadybos sistema, šis sertifikatas baigia galioti: 2021 m. Birželio 17 d

Sertifikato Nr .: DK009654 Versija: 1 Patikrinimo data: 2018 m. Birželio 12 d

MIX responsible Paper fromsources FSC FSC^{C} C074440

Certification Office: Bureau Veritas Certification Denmark A/S - Oldenborggade 25-31, 7000 Fredericia, Denmark

Further clarifications regarding the scope of this certificate and the applicability of the Management System requirements may be obtained by consulting the DANAK organization. To check this certificate validity, please call (+45) 77 311 000. SYSTEM Reg.nr. 5005