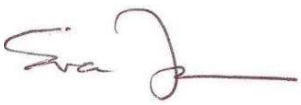
EC Declaration of Conformity

Manufacturer:	AKLA AB
Manufacturers Address:	(Enhagsslingan 2, SE-187 40 Täby), Box 534, SE-182 15 Danderyd, Sweden
SRN (Single Registration Number):	Not applicable
Authorized Representative Name (if applicable):	Not applicable
Authorized Representative Address (if applicable):	Not applicable
Basic UDI-DI:	Not applicable
Name of the Device (s):	Wound swab, IVF HARTMANN AG
Product code:	94408(HA)
Classification:	Class I, Rule 1
Notified Body name:	Not applicable
Notified Body Address:	Not applicable
Notified Body Identification number:	Not applicable

This declaration of conformity is issued under the sole responsibility of AKLA AB.
We hereby declare that he medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by the Quality System approval to ISO 13485 issued by Intertek Certification AB
All supporting documentation is retained at the premises of the manufacturer.

Signature:



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Eva Janmark
Managing director

Place and date (yyyy.mm.dd) of issue:

Täby 2020-06-01

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