



EC CERTIFICATE

According to Annex II (excluding article 4) of the Directive 93/42/EEC on Medical Devices

Full Quality Assurance System

Certificate Number

CE-MDD-0021/01/2020/01

Manufacturer : [REDACTED]
Address : [REDACTED]
Country : [REDACTED]

Product(s) : Implant, Hyaluronic Acid Gel with Lidocaine

Model(s) : Please see 2nd page for details of product

Report Number : CE-MDD-0021

First Issue Date and Place : 2020.01.21 & Istanbul

Revision Number and Date : 00 / --

Validity Date : 2024.05.26

NOTICE, Notified Body 2764, declares that the aforementioned manufacturer has implemented a quality assurance system according to Annex II, Section 3 of the directive 93/42/EEC on medical devices. This quality assurance system covers those aspects of manufacturing concerned with securing and maintaining safe conditions of the respective product(s) and conforms to the provisions of this Directive. The approved quality system is subject to surveillance pursuant to Annex II, Section 5 of Directive 93/42/EEC and unannounced audits. NOTICE must be informed of any significant changes in the design and/or construction of the product(s). This certificate is valid together with Design Certificate issued according to 93/42/EEC Annex II (Section 4) for products defined below.

Özlem Vicdan Akdağ
Chairman



**NOTİCE BELGELENDİRME
MUAYENE VE DENETİM
HİZMETLERİ A.Ş.**

Esentepe Mahallesi Milangaz Caddesi
No:75/A/92 Kartal / İstanbul / TÜRKİYE
www.notice.com.tr



Trade Name	Model (s) / Product code	Volume (ml)
[REDACTED]	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
[REDACTED]	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
[REDACTED]	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
STARFILL	Implant Plus Lidocaine	1.1
	Deep Plus Lidocaine	1.1
	Plus Lidocaine	1.1
[REDACTED]	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
[REDACTED]	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]
	[REDACTED]	[REDACTED]



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