EC Certificate
Full Quality Assurance System

Certificate No.:	Project No.:	Valid Until:
11299-2017-CE-IND-NA-PS Rev 0.0	**PRJC-217463-2010-PRC-IND**	**26 July 2020**

This is to certify that the quality system of:

Kehr Surgical Private Limited
C-34, Panki Industrial Estate,
Kanpur – 208 022, U.P., India

For design, production and final product inspection/testing of:

Sterile Disposable Medical Devices

Has been assessed with respect to:

The conformity assessment procedure described in Article 11.3.a and Annex II excluding section 4 (Module H2) of Council Directive 93/42/EEC on Medical Devices, as amended

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and Date:
Høvik, 19 October 2017

NORWEGIAN ACCREDITATION
PROD 021
Notified Body No.: 2460

For:
DNV GL NEMKO PRESAFE AS

Alessandra Rinna

The Certificate has been digitally signed.
See www.presafe.com/digital_signatures for more info

Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.

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Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as “Forskrift for Medisinsk Utstyr” by the Norwegian Ministry of Health and Care Services.

Certificate history

Revision	Description	Issue Date
0.0	Supersedes DNVGL (NB 0434) Certificate no: 81770-2010-CE-IND-NA 1.0 following transfer to notified body functions to DNV GL Nemko Presafe AS (NB 2460)	2017-10-19

Products covered by this Certificate

Product Description	Product	Class
Surgical Blades (Sterile & Non Sterile) in Carbon Steel & Stainless Steel with or without Twin Rib	Sizes: 9, 10, 10A, 11, 11P, 12, 12B, 12D, 13, 14, 15, 15A, 15B, 15C, 15T, 16, 17, 18, 19, 20, 21, 22, 22A, 23, 24, 24D, 25, 25A, 26, 27, 34, 36, 36D, 60, PM40, PM40B, PM60, PM60B	IIa
Gouge Blades (Sterile & Non Sterile) in Carbon Steel & Stainless Steel	1, 1V, 2, 2M, 2V, 3, 3V, 4, 5, 6, 8, 10, 12, 15	IIa
Disposable Scalpels (Sterile & Non Sterile) in Carbon Steel & Stainless Steel with or without Twin Rib	Sizes: 9, 10, 10A, 11, 11P, 12, 12B, 12D, 13, 14, 15, 15A, 15B, 15C, 15T, 16, 17, 18, 19, 20, 21, 22, 22A, 23, 24, 24D, 25, 25A, 26, 27, 34, 36, 36D, 60, PM40, PM40B, PM60, PM60B, Mini Stitch Cutter	IIa
Disposable Safety Scalpels (Sterile & Non Sterile) in Carbon Steel & Stainless Steel with or without Twin Rib	Sizes: 9, 10, 10A, 11, 11P, 12, 12B, 12D, 13, 14, 15, 15A, 15B, 15C, 15T, 16, 17, 18, 19, 20, 21, 22, 22A, 23, 24, 24D, 25, 25A, 26, 27, 34, 36, 36D, 60, PM40, PM40B, PM60, PM60B, Mini Stitch Cutter	IIa
Thumb Scalpel (Sterile & Non Sterile) in Carbon Steel & Stainless Steel with or without Twin Rib	9, 10, 10A, 11, 11P, 12, 12B, 12D, 13, 14, 15, 15A, 15B, 15C, 15T, 16, 17, Mini Stitch Cutters	IIa
Stitch Cutter Blades (Sterile & Non Sterile) in Carbon Steel & Stainless Steel with or without Twin Rib	Long, Standard and Mini	IIa

Podiatry, Miniature & Fine Range of Surgical Blades (Sterile & Non Sterile) in Carbon Steel & Stainless Steel	61, 61S, 62, 62SB, 63, 64, 65, 65A, 67, 68, 69, 90, 91, 312, 313, 314, 316	IIa
Grafting Blades (Sterile & Non Sterile) in Carbon Steel & Stainless Steel	Skin Graft, Microtome Blade	IIa

The complete list of devices is filed with the Notified Body.

Sites covered by this certificate

Site Name	Address
Kehr Surgical Private Limited	C-34, Panki Industrial Estate, Kanpur – 208 022, U.P., India

EU Representative

Medical Device Safety Service, Schiffgraben 41, 30175, Hannover, Germany.

Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

End of Certificate