CERTIFICATE * СВИДЕТЕЛЬСТВО * ZERTIFIKAT * CERTIFICATO

Certificate of Compliance

We hereby declare that the technical file of product complied with the requirement of Medical Devices Directive 93/42/EEC

INDUS MEDICARE PRIVATE LIMITED

Registered Office: 1-10-60/3, "Suryodaya", Begumpet, Hyderabad, Telangana State, India- 500 016

Factory: Survey No 129(Part), NH 44, Ramalpally, Manoharabad Mandal, Medak District, Telagana State, India- 503 226

Product:

Manufacturing & Supply of Water Based Personal Lubricant

The certification body has performed an audit of the above product testing documents & verified the design system, manufacturer and final inspection of the certified product. The product specification has been approved and is subject to continuous surveillance according to the Directive requirement.

In accordance with MDD 93/42/EEC, its hereby declared

The manufacturer has produced evidence that he fulfills the requirements of the standard Medical Devices Directive 93/42/EEC for execution of above mentioned equipment referring to the intended use, the certification body has conducted with successful results the review of the Manufacturer's technical documents of the certified products.

This Certificate is issued under the following condition

1- The Manufacturers technical documentation as required has been reviewed and found to comply with the requirements of standard

2- The certificate remains valid until the manufacturing condition are changed

3- The certificate validity is conditioned by positive results or surveillance audit

4- Any significant changes in the design or process use to manufacture the product or revision to the directive or standards referred above may require special audit.

5- This certificate will remain valid till test reports of the products are in line with Medical Devices Directive 93/42/EEC standard

6- After fulfilling the relevant standard testing performance, the manufacturer shall affix to each device, of the referenced model

The compliance as shown above can be used , under the responsibility of the manufacturer, after completion of a declaration of conformity and compliance with all relevant standard requirements. The statement is based on a single evaluation of one sample of above mentioned product it does not imply an assessment of the whole production

Signed for and behalf of QVR

Certificate Number: CE70XXII120208440

This certificate is valid from 22-July-2021 until 21-July-2024 and remains valid subject to satisfactory surveillance audits on or before 21-July-2022 and 21-July-2023

Re-certification audit due on 21- July-2024

Directorate of Accreditation for Assessment Services

973 Elizabeth St. Strathmore Alberta - T0C Canada

www.daasaccreditation.org / www.qvrcerts.com

Please Visit to verify the validity of this certificate on www.daasaccreditation.org or scan QR code

To verify the validity of this unaccredited certificate please visit www.daasaccreditation.org or scan QR code. Surveillance audit [illegible] year. This is to certify that the management system of this company has been [illegible] the certified client does not allow surveillance, re-certification audits, certificate would be re[illegible]. This certificate remains the property of Directorate of Accreditation for Assessment Services (DAAS). [illegible] certificate is Recognized by Directorate of Accreditation for Assessment Services (DAAS).

Digitally signed by Chirtoacă Iurie
Date: 2023.01.03 13:21:32 EET
Reason: MoldSign Signature
Location: Moldova