CERTIFICATE * СВИДЕТЕЛЬСТВО * ZERTIFIKAT *

Certificate of Conformity

We hereby declare that the technical file of product complied with the requirement of Directive (EU) 2019/904

CHETAK MEDIPLAST INDUSTRIES

49/2/1 AND 49/2/1/1, MS LAYOUT, SRIGANDAKAVAL, SUNKADAKATTE, BANGALURU (BANGALORE) URBAN, KARNATAKA, 560091, INDIA.

Product:
Annexure 1 to Annexure 2

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 Directive (EU) 2019/904, its hereby declared
The manufacturer has produced evidence that he fulfills the requirements of the standard Directive (EU) 2019/904 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 Directive (EU) 2019/904 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: DAAS-CMI-000777

This certificate is valid from 01-July-2025 until 30-June-2028
Ist Surveillance audit due date 30-June-2026
IInd Surveillance audit due date 30-June-2027
Re-certification due date 30-June-2028

Directorate of Accreditation for Assessment Services
973 Elizabeth St. Strathmore Alberta - T0C Canada
www.daasaccreditation.org

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 shall be conducted at least once a calendar year. This is to certify that the management system of this company has been found to confirm to the above. If the certified client does not allow surveillance, re-certification audits, certificate would be return to Quality Verification Registrar. This certificate remain the property of Directorate of Accreditation for Assessment Services (DAAS). and this certificate is Recognized by Directorate of Accreditation for Assessment Services (DAAS).

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CHETAK MEDIPLAST INDUSTRIES

49/2/1 and 49/2/1/1, MS Layout, Srigandakaval, Sunkadakatte,
Bangaluru (Bangalore) Urban, Karnataka, 560091, India.

Annexure: 1

SL. No.	Products
1	Bedpan Plastic
2	Beakers
3	Commode Pan And With Urinal Pot
4	Dispensing Cups
5	Enema Can Set
6	Eye Wash Cup
7	Face Mask (Particular Respirator)
8	Glycerin Syringe
9	Incentive Spirometer
10	Jalneti Pot
11	Kidney Tray Plastic

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49/2/1 and 49/2/1/1, MS Layout, Srigandakaval, Sunkadakatte, Bangaluru (Bangalore) Urban, Karnataka, 560091, India.

Annexure: 2

SL. No.	Products
12	Measuring And Scientific Jars
13	Sputam Cups
14	Sitz Bath
15	Sharp Conatiner
16	Urine Sample Containers
17	Urine Can Plastic (Hand-Held Urinal)
18	Vaporizer (Steam Inhaler)

Signed for and behalf of QVR

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