



CIN : L22193DL2024PLC428183

ANONDITA MEDICARE LIMITED

AN ISO 9001:2015 ISO 13485:2016 & CDSO CERTIFIED CO.
Manufacturer of Condoms

February 11, 2026

To,
The Manager,
Listing Compliance Department
National Stock Exchange of India Limited
Exchange Plaza, Bandra Kurla Complex,
Bandra (East), Mumbai-400051

Sub: Disclosure under Regulation 30 of the Securities and Exchange Board of India (Listing Obligations and Disclosure Requirements) Regulations, 2015

Ref: Disclosure under Regulation 30 dated January 10, 2026 regarding receipt of MDSAP Certification

SYMBOL: ANONDITA
ISIN: INE0VTV01012

Dear Sir/Madam,

Pursuant to Regulation 30 read with Sub-paragraph 12 of Paragraph B, Part A of Schedule III of the SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 ("Listing Regulations"), and in continuation of the Company's disclosure dated January 10, 2026 intimating the receipt of the Medical Device Single Audit Program (MDSAP) Certification, we hereby inform that the Company has received the **70951 Medical Device Single Audit Program (MDSAP) Certification vide OPINION No. 0090436/26-6** for Good Manufacturing Practices for Medical Devices from the Industry International Except Mercosur, with Brazil in scope, issued by DQS Medizinprodukte GmbH, an accredited auditing body **and recognized by ANVISA – National Health Surveillance Agency, (Brazil)**, valid for a period of four (4) years.

This certification is expected to enhance the Company's regulatory compliance and market access in Brazilian market.

The details as required under Reg. 30 of the SEBI (Listing Obligations and Disclosure Requirements), Regulations 2015 read with Para B of Part A to Schedule III and SEBI Master Circular No. HO/49/14/14(7)2025-CFD-POD2/I/3762/2026 dated 30th January 2026, are also enclosed herewith as Annexure-A.

The said information will also be uploaded on the website of the company <https://anonditamedicare.com/>.

Kindly take the above information on your record.

Thanking You.
Yours faithfully,

For and on behalf of
ANONDITA MEDICARE LIMITED

Bhawna Bisht
Company Secretary and Compliance Officer
M. No. A70843



Regd. Off.: Flat No. 704, Narmada Block N-6, Sector - D, Block - C, Vasant Kunj, New Delhi -110070
Corp Off. : D-001, Sector-80, Noida-201305, (U.P.) INDIA
Tel.: 0120-4520300/1/2/3 till 99 (100 Lines) Fax : 0120-4520314
E- mail : info@anonditamedicare.com | accounts@anonditahealthcare.com
Website: www.anonditamedicare.com



ANONDITA MEDICARE LIMITED

AN ISO 9001:2015 ISO 13485:2016 & CDSCO CERTIFIED CO.
Manufacturer of Condoms

Annexure A

Disclosure of information pursuant to Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulations 2015 read with Schedule III and SEBI Master Circular No. HO/49/14/14(7)2025-CFD-POD2/I/3762/2026 dated 30th January 2026.

Name of the regulatory or licensing authority	ANVISA – National Health Surveillance Agency (Brazil) DQS MedizinprodukteGmbH (Auditing/Certification Body)
Brief details of the approval/license obtained/withdrawn/surrendered	Grant of “ 70951 – MDSAP – Medical Devices – Certification of Good Manufacturing Practices for Medical Devices from the Industry International Except Mercosur ”, with Brazil (ANVISA – National Health Surveillance Agency) in scope. The certification confirms that the Company’s manufacturing facility (Anondita Medicare Limited) has been recognized for Good Manufacturing Practices (GMP) for Health Products – Class III Medical Devices for the Brazilian market. Applicant/Distributor in Brazil: Star Pharma Importadora e Exportadora Ltda (pursuant to the Distributor Agreement executed between the Applicant and the Company dated June 16, 2025) Manufacturer: Anondita Medicare Limited Operating Permit: 8223411 Certificate: Good Manufacturing Practices for Health Products – Class III Medical Devices The approval is supported by information from regulatory authorities and third-party auditing bodies recognized by ANVISA.
Impact/relevance of such approval/license to the listed entity;	As the Company is engaged in the manufacturing of male and female condoms, the grant of the Medical Device Single Audit Program (MDSAP) Certification, recognized by ANVISA – National Health Surveillance Agency, Brazil, enables the Company to sell, distribute, and market its products in Brazil and also enhances the Company’s regulatory credibility in this jurisdictions.
Withdrawal/cancellation or suspension of licence/approval by the regulatory or licensing authority, with reasons for such action, estimated impact (monetary or otherwise) on the listed entity and penalty, if any	Not applicable.
Subsequently, the listed entity shall inform the stock exchange(s), the actual impact (monetary or otherwise)	Not applicable.





CIN : L22193DL2024PLC428183

ANONDITA MEDICARE LIMITED

AN ISO 9001:2015 ISO 13485:2016 & CDSCO CERTIFIED CO.
Manufacturer of Condoms

along with corrective actions taken by the listed entity pursuant to the withdrawal, cancellation or suspension of the key license/ approval.



UNIT



OFFICE



DIPPING AREA



TESTING AREA

Regd. Off.: Flat No. 704, Narmada Block N-6, Sector - D, Block - C, Vasant Kunj, New Delhi -110070
Corp Off. : D-001, Sector-80, Noida-201305, (U.P.) INDIA
Tel.: 0120-4520300/1/2/3 till 99 (100 Lines) Fax : 0120-4520314
E- mail : info@anonditamedicare.com | accounts@anonditahealthcare.com
Website: www.anonditamedicare.com