CIN : L22193DL2024PLC428183
ANONDITA MEDICARE LIMITED
AN ISO 9001:2015 ISO 13485:2016 & CDSCO CERTIFIED CO.
Manufacturer of Condoms

January 10, 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.: SEBI Circular: SEBI/HO/CFD/PoD2/CIR/P/0155 dated 11th November 2024

SYMBOL: ANONDITA
ISIN: INE0VTV01012

Dear Sir/Madam,

Pursuant to the provisions of 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"), we hereby inform that the Company has received the Medical Device Single Audit Program (MDSAP) Certification in accordance with ISO 13485:2016 from **DQS Medizinprodukte GmbH**, a designated certification authority.

The receipt of this certification enables the Company to market its products in five international regulatory jurisdictions, namely Australia, Brazil, Canada, Japan, and the United States of America.

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 circular SEBI/HO/CFD/PoD2/CIR/P/0155 dated 11th November 2024, 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
Digitally signed by Bhawna Bisht
Date: 2026.01.10 16:49:45 +05'30'

Bhawna Bisht
Company Secretary and compliance officer
M. No. A70843

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Annexure A

Details as required under SEBI (Listing Obligations and Disclosure Requirements) Regulations, 2015 read with SEBI Circular No. SEBI/HO/CFD/PoD2/CIR/P/0155 dated 11th November 2024.

Name of the regulatory or licensing authority	DQS Medizinprodukte GmbH
Brief details of the ~~approval~~/license ~~obtained/ withdrawn/ surrendered~~	Grant of **MDSAP Certification (ISO 13485:2016)** confirming compliance with Quality Management System requirements for medical device manufacturing, and the certification enhances regulatory credibility of the Company, enables acceptance of audits across multiple international regulatory jurisdictions, strengthens compliance framework, and supports business expansion in regulated medical device markets.
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 enables the Company to sell, distribute, and market its products in five major international regulatory jurisdictions, namely Australia, Brazil, Canada, Japan, and the United States of America, and also enhances the Company's regulatory credibility in these 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) along with corrective actions taken by the listed entity pursuant to the withdrawal, cancellation or suspension of the key license/ approval.	Not applicable.