

EU Declaration of Conformity

Name and address of the manufacturer:

Dr. Arabin GmbH & Co. KG, Alfred-Herrhausen-Str. 44, 58455 Witten, Germany

We declare under our sole responsibility that the medical device

| Product name | Article Code / REF Nr. | Class |
|-----------------------------------|-------------------------|-------|
| Cube Pessary | WP 0 - 9 | Ila |
| Cube Pessary perforated | WPP 0 - 9 | Ila |
| Tandem Pessary | TP 0+0 F0 – TP 5+5 F5 | Ila |
| Tandem Pessary perforated | TPP 0+0 F0 – TPP 5+5 F5 | Ila |
| Cup Pessary (former Club Pessary) | KP050 - 090 | Ila |

Basic UDI-DI: 426024582GYNCP8Q

SRN: DE-MF-000005474

complies with the requirements of Regulation (EU) 2017/745 and other EU Guidelines that are applicable.

Indication of the risk class and the conformity assessment procedure:

Class Ila in accordance with Annex VIII of Regulation (EU) 2017/745

Conformity assessment procedure according to Annex IX Chapters I + III

Name and identification number of the Notified Body:

DNV MEDCERT GmbH

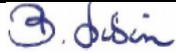
Identification number: 0482

Certificate with validity:

Certificate ID: 1340GB448241217

Valid until: 2028-07-02

This declaration of conformity is valid until the above certificate expires or if a change is made to the product.

| Department | Date: | Released by: |
|--|------------|---|
| CEO / PRRC – Prof. Dr.med. Dr.h.c.mult. B.Arabin | 05.01.2026 |  |
| QM Representative – M. Prior | 05.01.2026 |  |