

Declaration of Conformity

Product identification

Product name: KY Jelly (sterile)
Brand: KY/K-Y/Durex K-Y/ KY Durex/ K-Y Durex/ Durex
Reference: MDTF 198
Issue Number: 06
Component number: 8203148

Manufacturer

Name: RB Healthcare (UK) Ltd
Address: Dansom Lane, Hull, HU8 7DS
Country: United Kingdom

Authorized Representative/Distributor in Europe

Name: RB NL Brands B.V
Address: Schiphol Boulevard 207, 1118 BH Schiphol.
Country: Netherlands

Registration Information

Notified Body ID: 1639
Name: SGS Belgium NV
Address: Noorderlaan 87, BE-2030 Antwerpen.
Country: Belgium

CE Certificate No.: GB19/963047
Date CE marked: 8th April 2019
Originally CE Marked 22nd February 2016 under SGS UK, 0120. CE Certificate number GB99/50557.01.

Classification: Class: IIa Rule: 5
Route to compliance: Annex II (Excluding section 4) of MDD 93/42/EEC Council Directive (as amended)
Standard Applied: MDTF 198

Reckitt Benckiser Healthcare (UK) declares that the above-mentioned product is in conformity with the essential requirements and provisions of Council Directive 93/42/EEC as amended and conforms to the standards listed in MDTF 198.

Signatures

Name and function:

Hannah Hussey

Global Regulatory Manager – Intimate
Wellness Compliance Manager

Signature:

 Electronically signed by: Hannah Hussey
Reason: I approve this document.
Date: Sep 9, 2022 07:15 GMT+1

Date:

09-Sep-2022

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Final Audit Report

2022-09-09

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-  Document created by Maryam Roostan (Maryam.Roostan@reckitt.com)
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Date (GMT)	Signed by	Reason
14-Sep-2022 12:14:05	Brookes Rebecca (RBrookes)	Signing as Approver