EU DECLARATION OF CONFORMITY

DOC NAME						PAGES
EU Declaration of Conformity MDR - Navina Consumable Set						
DOC TYPE	DOC NO	VERSION	ENCLOSURE	DATE	STATE	
DC	10104	В	None	2023-12-19	Approved	
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We,

Wellspect HealthCare Aminogatan 1, P.O. Box 14, SE-431 21 Mölndal, Sweden

being the Manufacturer, registered in the European Union under Single Registration Number (SRN) No. SE-MF-000026271, of the Navina System Consumable Set, including the products listed in the Annex I to this document, with the following characteristics:

- device class IIa, as determined by Rule 5, according to Regulation (EU) 2017/745, Annex VIII
- intended for Transanal irrigation
- GMDN code: 61094
- EMDN category G / code(s):
 - o G020102 Intestinal Lavage tubes and sets
- Basic UDIDI/Global Model Number: 733338724106D7

Declare under our sole responsibility that the product(s) conform to the requirements of Regulation (EU) 2017/745 of the European Parliament and of the Council, on medical devices, and meet(s) the relevant General Safety and Performance Requirements of Annex I

All devices are designed, manufactured, tested, and released for sale in accordance with the technical documentation according to Annex II and III of regulation (EU) 2017/745 and the applicable standards.

The conformity assessment procedure was performed following Annex IX of EU Regulation 2017/745.

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This declaration is made based on the EU Technical Documentation Assessment Certificate MDR 780135 issued by the Notified Body:

BSI Group the Netherlands B.V. (2797) Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, Netherlands

This declaration of Conformity is approved and signed as dated on first page.

Wellspect HealthCare, Mölndal, Sweden

TONI JORGENSEN

VICE PRESIDENT QUALITY ASSURANCE & REGULATORY AFFAIRS

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ANNEX I

Article (model) No.*	Product Name, Description		
69003	Navina Consumable Set Regular		
69004	Navina Consumable Set Small		
69016	Navina Consumable Set Cone		

^{*}Generic article number without the 2-digit suffix specific for a region or country destination when distributing an article.

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REVISION HISTORY

Document Version	Change note/Description
В	New Basic UDI-DI assigned due to an error in previous Basic UDI-DI identifier.