

	Technical File – Declaration of Conformity	WI-EU-REG-040 – Annex IV	
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We

Name + address + Single Registration Number of manufacturer:	Laboratories Anios 1 RUE DE L'ESPOIR 59260 LEZENNES FRANCE
Name + address + Single Registration Number of authorized representative:	Not Applicable



declare on our sole responsibility that

the medical device	ANIOS LB 400
Type / Intended Use	Detergent
Class Rule according to MDR Annex VIII	I Rule 1

Meets all the provisions of the Regulation (EU) 2017/745 on medical devices.

Notified body name, address, ID	Not Applicable
Conformity assessment procedure	Article 52, MDR (Annex II + III)
ID of the certificates issued	Not Applicable
Common Specifications	Not Applicable
Validity	26.05.2024

Valid in conjunction with the batch related release documentation

Monheim, 20 August 2021	Dieter Wirbals 	Sainghin, 20 August 2021	Jean-Noël Bertho 
Place, date	Name and function (Person Responsible for Regulatory Compliance)	Place, date	Name and function (RD&E Representative)

Signed on behalf of: *[Laboratoires Anios]*

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Medical Device	Brand	Device Subcategory	Part Number / SKU	Basic UDI-DI
Anios LB 400	Anios	Detergent	21610381X	359761LA01C150INQ3
Anios LB 400	Anios	Detergent	21610383T	