LABORATOIRES ANIOS An Ecolab Company	TECHNICAL FILE - EU DECLARATION OF CONFORMITY (MDR)	**Réf : FQ279B** **Date : 21/07/22**

We

Name + address + Single Registration Number of manufacturer:	**Laboratoires Anios** **1 rue de l'Espoir** **59260 Lezennes** **France** **FR-MF-000011756**
Name + address + Single Registration Number of authorized representative:	Not Applicable

declare on our sole responsibility that

the medical device	ANIOSYME FOAM
Type / Intended Use	*DETERGENT INVASIVE / Instrument cleaning*
Class	*I*
Rule according to MDR Annex VIII	*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	*MDR (EU) 2017/745 article 52, Annex II and III*
ID of the certificates issued	Not Applicable
Common Specifications	Not Applicable
Validity	26 May 2024

Valid in conjunction with the batch related release documentation

Sainghin 09/01/2023	Digitally signed by Dieter Wirbals DN: cn=Dieter Wirbals, o=Medical Device, ou=Regulatory Affairs, email=dieter.wirbals@ecolab.com, c=DE Date: 2023.01.12 14:54:42 +01'00' Dieter Wirbals	Sainghin 09/01/2023	Digitally signed by Jean-Noël BERTHO Date: 2023.01.13 17:45:58 +01'00' 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**

Medical Device	Brand	Device Subcategory	Packaging variant	Basic UDI-DI
ANIOSYME FOAM	ANIOS	Detergent invasive	544 12x750mL	359761LA0150C1LIME