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	ANIOS LB 30
Type / Intended Use	*RINSING AGENT INVASIVE / Acid rinsing for thermal wash basins*
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 09/01/2022	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 14:52:40 +01'00' Dieter Wirbals	Sainghin 09/01/2022	Digitally signed by Jean-Noël BERTHO Date: 2023.01.13 17:35:54 +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
ANIOS LB 30	ANIOS	Rinsing agent invasive	38 2x5L	359761LA01501NLIN3