

	Technical File – Declaration of Conformity	WI-EU-REG-040 – Annex IV	
		Revision: 01	Page 1 of 2

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 RDA
Type / Intended Use	Rinsing Agent
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]*

Medical Device	Brand	Device Subcategory	Part Number / SKU	Basic UDI-DI
Anios RDA	Anios	Rising Agent	2372018FA	359761LA01C150INQ3
Anios RDA	Anios	Rising Agent	2372038HD	
Anios RDA	Anios	Rising Agent	2372038UF	
Anios RDA	Anios	Rising Agent	2372015UG	
Anios RDA	Anios	Rising Agent	2372018UG	
Anios RDA	Anios	Rising Agent	2372271UG	
Anios RDA	Anios	Rising Agent	2372331UG	
Anios RDA	Anios	Rising Agent	2372024UG	
Anios RDA	Anios	Rising Agent	2372038UG	
Anios RDA	Anios	Rising Agent	2372024HD	
Anios RDA	Anios	Rising Agent	2372038UD	
Anios RDA	Anios	Rising Agent	23720153T	
Anios RDA	Anios	Rising Agent	2372038UE	