

Declaration of Conformity

Product Name	Cryovial™			
Product Codes	T210; T301; T308; T309; T310; T311 Series			
Manufacturer	Simport Scientific Inc. 2588 Bernard-Pilon Beloil Quebec J3G 4S5 Canada			
Authorized Representative in Europe	<table border="1" style="display: inline-table; vertical-align: middle;"> <tr> <td style="text-align: center; width: 50px; height: 50px;">EC</td> <td style="text-align: center; width: 50px; height: 50px;">REP</td> </tr> </table>	EC	REP	EMERGO EUROPE Prinsessegracht 20 2514 AP The Hague The Netherlands
EC	REP			

Simport declares that the above-mentioned products meet the provision of the Council Directive 98/79/EC for *In Vitro* Diagnostic Medical Devices and Directive 98/79/EC as transposed in the national laws of the Member States:

- That the products have been classified as general IVD and class A (Low Individual Risk and Low Public Health Risk);
- That the products listed below are in conformity with the Annex III (Sections 1 to 5), and annex I (Essential Requirements) of Directive 98/79/CE;
- That the products do not contain medicinal substances;
- That the products do not contain animal tissues.



Annette Roy, Regulatory and Technical Support
Place and Date: Beloil, May 03, 2022

Addendum to the original Declaration of Conformity

Per 31 January 2023, the address of our EU Authorized Representative as listed on the original DoC has changed.

Old address AUTHORIZED Representative		
Name of company	Address	Telephone/email
Emergo Europe	Prinsessegracht 20 2514 AP The Hague The Netherlands	+31.70.345.8570 - phone EmergoEurope@ul.com

New address AUTHORIZED Representative		
Name of company	Address	Telephone/email
Emergo Europe	Westervoortsedijk 60 6827 AT Arnhem The Netherlands	+31.70.345.8570 - phone EmergoEurope@ul.com

DATE: 01/03/2023

COMPANY REPRESENTATIVE: CLAUDE LEBOND
 TITLE: VICE-PRESIDENT SIGNATURE: [Signature]

DATE: 19/09/2024