



# Declaration of Conformity

Transport Vials Range (Filled & Un-filled)

		Doc No.	TD003
Declaration of Conformity		Version. No.	01
		Issue Date	10 May 2022
Prepared By	Sajana Vattikuti	Pages	1 of 4
Approved By	Janet Mackenzie		



**EC Declaration of Conformity**  
**European Communities Council Regulation EU 2017/746**  
**Concerning IVD Medical Devices**

**Manufacturer:** **APACOR LIMITED**  
**Unit 5, Sapphire Centre**  
**Fishponds Road**  
**Wokingham**  
**Berkshire RG41 2QL**  
**United Kingdom**

**EC Authorised Representative :** **Medical Device Safety Service GmbH (MDSS)**  
**Schiffgraben 41**  
**30175 Hannover**  
**Germany**

**SRN:** **DE-AR-000005430**

We hereby declare that the following mentioned products meet the provisions of the Council Regulation EU 2017/746 covering medical devices. All documentation is controlled and retained on company premises. This declaration of conformity is issued under the sole responsibility of Apacor Limited.

**Product:** See list enclosed

**Classification:** **Class A RULE 5 (C )**

**GMDN :** **63816**

**Standards Applied:**

Document Reference	Description
BS EN ISO 14971:2019	Medical devices – Application of risk management to medical devices
BS EN ISO 18113-2:2011	In Vitro Diagnostic Medical Devices - Information Supplied by the Manufacturer (Labelling) - Part 2: In Vitro Diagnostic Reagents for Professional Use
ISO 13485:2016	Medical device – Quality management system – Requirements for regulatory purposes
BS EN ISO 15223-1:2021	Medical Devices -- Symbols to be Used with Medical Device Labels, Labelling and Information to be Supplied -- Part 1: General Requirements
BS EN ISO 13612:2002	Performance Evaluation of In Vitro Diagnostic Medical Devices
BS EN ISO 23640:2015	In vitro diagnostic medical devices — Evaluation of stability of in vitro diagnostic reagents
MEDDEV 2.12-1 rev. 8,2013	Guidelines on a medical devices vigilance system

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**Notified Body** N/a

**Date of Issue:** 10 May 2022

**Approved By:**

Sajana Vattikuti

QA/RA Manager

Apacor Ltd

### Change Control History

Version	Change Description	Reason for change	Signed
01	First Issue of Document	Requirement of EN ISO 13485:2016 and IVDR EU 2017/746	

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Product Description	Product Code	Pack Size
<b>30ml Transport Vial</b>		
No Additive	149970	40
15ml Formalin & Triton X	148998	40
15ml 10% Formalin & Triton X	151000	40
15ml Apafix™	160001	40
15ml SAF & Triton X	249400	40
15ml Alcorfix™	249420	40

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