

# Declaration of Conformity

## In accordance with the In Vitro Diagnostic Directive 98/79/EC

The following Devices, classified as “all other IVD Medical Devices”, conform to the relevant provisions of the IVD Directive 98/79/EC and are CE marked in accordance with Annex I and Annex III.

**Product:**

CEREBRO Sample Tracking System and associated accessories listed– See Appendix

**Manufacturer:**

Leica Biosystems Melbourne Pty Ltd  
495 Blackburn Road, Mt Waverley, Victoria 3149, Australia

**European Authorised Representative:**

CEpartner4U  
Esdoornlaan 13,  
3951 DB, Maarn,  
The Netherlands

Signed for and on behalf of:



Date: .....

9 Dec 2020

Adrienne Hardisty  
RA/QA Manager  
Leica Biosystems Melbourne Pty Ltd

## Appendix

The CEREBRO and associated accessories

<b><i>CEREBRO Software Product</i></b>	
Licence Key, Main, CEREBRO	65.6400
<b><i>CEREBRO Software Components/Accessories</i></b>	
Extended Licence Key - Accessioning Station	65.6410
Extended Licence Key - Grossing Station	65.6420
Extended Licence Key - Embedding Station	65.6430
Extended Licence Key - Sectioning Station	65.6440
Extended Licence Key - Send Out Station	65.6450
Extended Licence Key - Reporting Station	65.6460
Extended Licence Key - Processing Station	65.6461
Extended Licence Key - Staining Station	65.6462
Extended Licence Key - Universal Station	65.6463