

## Declaration of Conformity

|                                     |   |    |     |  |
|-------------------------------------|---|----|-----|--|
| Product Name                        | Histology Cassettes   |    |     |  |
| Intended use                        | Hold tissue specimens for further in vitro diagnostic   |    |     |  |
| Product Codes                       | M385; M386; M392; M405; M406; M407; M409; M460; M470; M471; M474; M475; M476; M478; M480; M482; M483; M485; M486; M490; M491; M492; M493; M498; M499; M502; M503; M505; M506; M507; M508; M509; M510; M511; M512; M515; M516; M517; M518; M520; M521; M525; M526; M430; M431; M432; M433; M434; M435; M530; M531 M532 M533 M534; M535 |    |     |  |
| Basic UDI-DI                        | 667243  |    |     |  |
| Manufacturer                        | Simport Scientific Inc.<br>2588 Bernard-Pilon<br>Beloeil<br>Quebec J3G 4S5<br>Canada  |    |     |  |
| Authorized Representative in Europe | <table border="1"> <tr> <td>EC</td> <td>REP</td> </tr> </table>   | EC | REP | <b>EMERGO EUROPE</b><br>Westervoortsedijk 60<br>6827 AT Arnhem<br>The Netherlands<br>SRN NL-AR-000000116 |
| EC                                  | REP   |    |     |  |

Simport declares that the above-mentioned products meet the provision of the Regulation (EU) 2017/746 of the European Parliament and of the Council on *In Vitro* Diagnostic Medical Devices and Regulation (EU) 2017/746 as transposed in the national laws of the Member States:

- That the products have been classified as general IVD and Rule 5(c), class A (Low Individual Risk and Low Public Health Risk);
- That the products listed above are in conformity with the Annex II (Essential Requirements) and III of Regulation (EU) 2017/746
- 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: Beloeil, February 02, 2023