

Declaration of Conformity

Product Name	Histosette I, Histosette II, Histosette II Quickload, Unisette, Unisette Quickload, Micromesh, Micromesh Quickload, , Slimsette, Slimsette Quickload, Microsette, Microsette Quickload, Swingsette, Swignsette Quickload, Macrosette, Superflow, Superflow Quickload, E-Z Load Histosette II, Cassettes in Quickload
Intended use	Hold tissue specimens for further in vitro diagnostic
Product Codes	M385; M386; M392; M393, M405; M406; M407; M409, M410, M430, M431, M432, M433, M434, M435, M460; M470; M471; M474; M475; M476; M477, 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; M525; M526;M530; M531; M532; M533; M534; M535;
Basic UDI-DI	667243
Manufacturer	Simport Scientific Inc. 2588 Bernard-Pilon Beloeil Quebec J3G 4S5 Canada
Authorized Representative in Europe	ECREPWestervoortsedijk 606827 AT ArnhemThe NetherlandsSRN NL-AR-000000116SRN NL-AR-000000116

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
- \rightarrow That the products do not contain medicinal substances;
- \rightarrow That the products do not contain animal tissues.

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Rossano Morgan-Cavallaro, QA/RA Agent Place and Date: Beloeil, May 28, 2024