KITAZATO®

EC DECLARATION OF CONFORMITY

Manufacturer:

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Dibimed

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Product: Trade Name: **Repro Plate**
Code number: **Refer to the following table.**

Code No.	Reference	Description
83006	Repro Plate-K1(6well)	6 well, 10 pieces in sterile pouch
83007	Repro Plate-K1(6well)	6 well, 1 piece in sterile pouch
83010	Repro Plate-K1(3well)	3 well, 10 pieces in sterile pouch
83011	Repro Plate-K1(3well)	3 well, 1 piece in sterile pouch
83016	Repro Plate-K2	6 well, 10 pieces in sterile pouch
83017	Repro Plate-K2	6 well, 1 piece in sterile pouch
83018	Repro Plate-K3	3 well, 10 pieces in sterile pouch
83019	Repro Plate-K3	3 well, 1 piece in sterile pouch
83020	Repro Plate-K6	6 well, 10 pieces in sterile pouch
83021	Repro Plate-K6	6 well, 1 piece in sterile pouch

Category: In vitro diagnostic medical devices
EDMS Code: 23.05 Histology / Cytology (HC) Culture Dishes

Conformity Assessment Route: **Annex III applied**

We herewith declare that the above-mentioned products meet the provisions of the following EC Council Directives and Standards. All supporting documentations are retained under the premises of the manufacturer.

General applicable directives: In vitro diagnostic medical devices: COUNCIL DIRECTIVE 98/79/EC (IVDD 98/79/EC)

Standards: Harmonized Standards (published in the Official Journal of the European Communities) applicable to this product are as per the "LIST OF APPLIED STANDARDS".

Signature: _____ Date: Dec. 18. 2014

Name: Futoshi Inoue
Position: President and Representative Director, Kitazato Corporation