

EC DECLARATION OF CONFORMITY

According to In vitro diagnostic Regulation (EU) 2017/746

Manufacturer: DLAB Scientific Co., Ltd.
YuAn Road 31, Airport Economic Core Zone, Shunyi District, Beijing 101318, China

SRN: CN-MF-000025396

European Representative: Kingsmead Service B.V.
Zonnehof 36, 2632 BE, Nootdorp, Netherland

SRN: NL-AR-000002066

Product Name: Low Speed Centrifuge

Product Model DM0412, DM0412 (A6-50P) , DM0506, DM0408, DM0424, DM0636, DM0306, Q0436E, Q0408E, DM0436E

Basic UDI-DI 697548995LD161AAUS

Classification: Class A, according to Rule 5a of IVDR Annex VIII

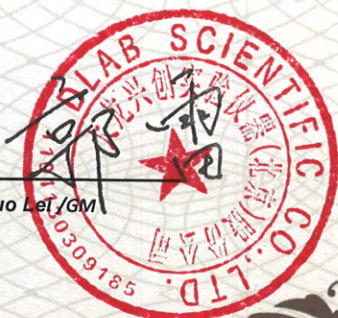
Applied Standards: EN ISO 13485:2016/A11:2021; EN ISO 15223-1:2021; EN ISO 14971:2019/A11:2021; EN ISO 18113-1:2011; EN 62366-1:2015; EN 61326-2-6:2006; EN 61010-2-101:2002

Conformity assessment procedure: Annex IV
According to Annex II and Annex III of TD

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the In vitro diagnostic Regulation (EU) 2017/746. All supporting documentations are retained under the premises of the manufacturer.

Signature: _____

Name/Position: Guo Lei / GM



Date: 2023/02/20

Place: Bei Jing / China