DECLARATION OF CONFORMITY ON MEDICAL DEVICE MANUFACTURER

20 July 2017
Vilnius

Medical device:	Dried blood spot sample collection card
Directive:	98/79/EB on *in vitro* diagnostic medical devices

„Bulleri group“, company code 301507397, hereby declares under its sole responsibility that the products RDA Card© dried blood spot sample collection cards necessary for blood samples collection and transportation to the laboratory according to Technical Regulation for the Safety of *in vitro* Diagnostic Medical Devices (Instruments) approved by Order No 679 of 29 December 2001 issued by the minister of Health of the Republic of Lithuania (hereinafter – Regulation) are classified as „other IVD medical devices” and are made in conformity with the applicable Essential Requirements on a given product specified in Annex I and Directive 98/79/EC of the European Parliament and of the Council on *in vitro* diagnostic medical devices.

Director Sigita Danilevičienė

IVD CE

ISO 13485:2003 CERTIFIED MATERIALS

ISO 9001:2008 CERTIFIED MATERIALS