PerkinElmer Singapore Pte Ltd
2 Tukang Innovation Grove
#04-01 JTC MedTech Hub @ MedTech Park
Singapore 618305

DECLARATION OF CONFORMITY

CE
Since
2013

The manufacturer **PerkinElmer Singapore Pte Ltd**

hereby declares that the **Laboratory Equipment (Class 8721-06 & 8721-86)**

of equipment models **WIZARD2**
2470-0010, 2470-0020, 2470-0050, 2470-0100, 2470-0150, 2470-0200, 2480-0010, 3470-0050, 3470-0100, 3470-0150, 3470-0200

specifically **Model No.:_____ Serial No.: _____ Mfg Date:_____**

conforms to the and its standards **EMC Directive (2014/30/EU)**
EN 55011:2009 + A1:2010 Group 1, Class A
EN 61326-1:2013 and EN 61326-2-6:2013
IEC 61000-4-2:2008; IEC 61000-4-3: 2010; IEC 61000-4-4:2004 + A1:2010;
IEC 61000-4-5:2005; IEC 61000-4-6:2008; IEC 61000-4-8:2009;IEC 61000-4-11:2004;
EN 61000-3-2:2006 + A1:2009 + A2:2009; EN 61000-3-3:2008)

as well as the and its standards **Low Voltage Safety Directive (2014/35/EU),**
EN 61010-1:2010
EN 61010-2-101:2017

IEC 61010-1:2010; IEC 61010-2-101:2015; CAN/CSA- C22.2 61010-1-12; CAN/CSA-C22.2 61010-2-101-15; UL 61010-1 (3rd Ed)

and the **In Vitro Diagnostic Directive IVDD 98/79/EC, Annex III**
Other device (all devices except Annex II and self-testing devices)

and also the **Restriction of Hazardous Substances in Electrical and Electronic Equipment Directive 2011/65/EU**
Beside an application exempted from the restriction in Article 4(1)
Specific to medical devices by Annex IV of RoHS Directive as below:
5. Lead in shielding for ionizing radiation.

Authorized Representative
EMERGO EUROPE
Prinsessegracht 20
2514 AP The Hague
The Netherlands

This declaration is issued under the responsibility of

Zhang Fu
as
Senior Manager Quality Assurance

PerkinElmer Document: 8845404, Rev G