PA 800 Plus, P/ACE MDQ Plus, CESI 8000 Plus

DECLARATION OF CONFORMITY

5302189

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(According to EN ISO/IEC 17050-1)

CE

We **AB SCIEX Pte. Ltd**

of **Blk 33, #04-06, Marsiling Ind. Estate Road 3, Woodlands Central Industrial Estate, Singapore 739256**

declare under sole responsibility that the product as originally delivered

Equipment: **Capillary Electrophoresis**
Product Name: **PA 800 Plus, P/ACE MDQ Plus, CESI 8000 Plus**

EU:

complies with the essential requirements of the following applicable European Directives, and carries the CE marking accordingly:

2006/42/EC	**- Machinery Directive**
2014/30/EU	**- Electromagnetic Compatibility Directive**
2012/19/EU	**- Waste Electrical and Electronic Equipment Directive**
2011/65/EU and 2015/863/EU	**- Restriction of Hazardous Substances Directive**

Australia and New Zealand:

complies with the requirements of the relevant ACMA Standards made under the *Radiocommunications Act 1992* and the *Telecommunications Act 1997 as well as* the New Zealand Radiocommunications Act 1989.

The product was tested in a typical configuration and conforms with the following standards:

Category	Standard		Classification
Safety	**EU:**	IEC 61010-1:2010+ A1: 2016 / EN 61010-1:2010 + A1:2019 IEC 61010-2-081:2019 / EN 61010-2-081:2020	Class 1
EMC	**EU:**	IEC 61326-1:2012 / EN 61326-1:2013	Class A
	Australia/New Zealand:	AS CISPR 11:2017	Class A
	Canada:	ICES-001 (CSA CISPR 11:2019)	Class A
	USA:	FCC Part 15	Class A

I, the undersigned, hereby declare that the equipment specified above conforms to the above Directives and Standards.

7/15/2022

X Nichole Riek

Signed by: Riek, Nichole

Nichole Riek
Vice President,
Quality Assurance and
Regulatory Affairs

EC Authorized Person
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