Biotage

Declaration of Conformity

Biotage Sweden AB, Box 8, SE-751 03 Uppsala, Sweden

Declare under our sole responsibility that the products

415000	TurboVap LV
415001	TurboVap II
415540	TurboVap EH

With S/N specified on the instrument in format: YYWWNNNNN
to which this declaration relates are in conformity with the following directives, standards and other normative documents

Directive 2014/30/EU	**Electro Magnetic Compatibility (EMC)** Standards used: EN 61326-1:2013 / IEC 61326-1:2012 CFR 47 FCC Part 15 Subpart B. Class A (2011)
Directive 2014/35/EU	**Low Voltage Directive** Standards used: EN/IEC 61010-1:2010 EN/IEC 61010-2-010:2014 EN/IEC 61010-2-081:2015
Directive 2012/19/EU	**WEEE** EN 50419:2006
Directive 2011/65/EC + 2015/863	**RoHS-2, -3** Standards used: SS-EN 50581:2012 SJ/T 11364-2014 (China RoHS)
For Canada and the USA	CAN/CSA-C22.2 No. 61010-1:2012-05 CAN/CSA-C22.2 No. 61010-2-010:2015-01 CAN/CSA-C22.2 No. 61010-2-081:2015-11 UL 61010-1:2012-05 UL 61010-2-010:2015-01 UL 61010-2-081:2015-11

Certificate No. U8 16 10 71091 022 issued by NCB TÜV SÜD America Inc.

Uppsala, 05 June 2019

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(Steve Jordan, Chief Scientific Officer - Biotage)