Declaration of Conformity

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

declare under our sole responsibility that the product

Biotage® PhyPrep™ **P/N: 416700**

to which this declaration relates are in conformity with the following directives, standards and other normative documents;

Directive 2006/42/EC Standards used:	**Machinery Directive (MD)** EN 61010-1:2010/A1:2019 EN 61010-2-081:2020
Directive 2014/30/EU Standards used:	**Electro Magnetic Compatibility (EMC)** EN 61326-1:2013 CFR 47 FCC part 15 Subpart B. Class A
Directive 2012/19/EU Standard used:	**WEEE** EN 50419:2006
Directive 2011/65/EU + 2015/863/EU Standard used:	**RoHS** EN 63000:2018
For Canada and the USA	UL 61010-1:2012 /R:2019-07 UL 61010-2-081:2019 IEC 61010-1:2010, IEC 61010-1:2010/AMD1:2016 IEC 61010-2-081: 2019 CAN/CSA-C22.2 No. 61010-1:2012/A1:2018-11 CAN/CSA-C22.2 No. 61010-2-081:2019 CFR 47 FCC part 15 Subpart B. Class A

NTRL Certificate No. U8 071091 0037 Rev. 00 issued by TÜV SÜD Product Service GmbH
CB Certificate No. DE 3 – 31447 issued by TÜV SÜD Product Service GmbH

Uppsala, 26 February 2021

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(Jon-Sverre Schanche, Chief Scientific Officer - Biotage)