Declaration of Conformity

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

declare under our sole responsibility that the product

Biotage® Extrahera™ **P/N: 414001**

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/IEC 61010-1:2010 IEC 61010-2-081/A1:2003
Directive 2014/30/EU Standards used:	**Electro Magnetic Compatibility (EMC)** EN/IEC 61326-1:2013 CFR 47 FCC part 15 Subpart B. Class A
Directive 2012/19/EU Standard used:	**WEEE** EN 50419:2006
For Canada and the USA	UL 61010-1:2012/R:2016-04 IEC 61010-1:2010 (CB) CAN/CSA-C22.2 No. 61010-1:2012/U2:2016-04 UL 61010-2-081:2015 IEC 61010-2-081:2015 (CB) CAN/CSA-C22.2 No. 61010-2-081:2015 CFR 47 FCC part 15 Subpart B. Class A Including national differences for Switzerland

NTRL Certificate No. U8 071091 0030 issued by TÜV SÜD Product Service GmbH
CB Certificate No. CB 071091 0029 issued by TÜV SÜD Product Service GmbH

Uppsala, 13 February 2019

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(Steve Jordan, Chief Scientific Officer - Biotage)