

Innovative Solutions for a Healthier World

Declaration of Conformity

Issued under the sole responsibility of:

Drucker Diagnostics 200 Shady Lane, Suite 170 Philipsburg, PA 16866 Tel: (814) 342-6205 Fax: (814) 342-4510

Object of Declaration:

Boost 2+ Flex, 00-783-243-000

This centrifuge is designed and manufactured in conformity with these EU directives:

- 2014/35/EU Low Voltage Directive
- 2014/30/EU Electromagnetic Compatibility (EMC) Directive
- 2011/65/EU RoHS Directive

This centrifuge is designed and manufactured in conformity with these standards:

Safety	UL61010-1/CSA C22.2 No. 61010-1-12 Third Edition: Safety Requirements for Electrical Equipment for Measurement, Control, and Laboratory Use, Rev. July 15, 2015
	IEC 61010-2-020:2006 / CSA C22.2 No. 61010-2-020-09 (R2014), Second Edition: Particular Requirements for Laboratory Centrifuges
EMC	FCC Part 15, Subpart B & ICES-003; IEC/EN 61326-1:2013 Electrical equipment for measurement, control and laboratory use. EMC requirements. Part 1: General requirements; IEC/EN 55011:2009 Industrial, scientific and medical equipment – Radio-frequency disturbance characteristics – Limits and methods of measurement. Also in conformity with these specifications: FCC Part 15, Subpart B / ICES-003 / EN 55011:2009 (Class A 150kHz-1GHz); EN 61000-3-2:2014; EN 61000-3-3:2013; EN 61000-4-2:2008; EN 61000-4-3:2010; EN 61000-4-4:2012; EN 61000-4-5:2014; EN 61000-4-6:2013; EN 61000-4-8:2009; EN 61000-4-11:2010

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10 November 2023

Signed for Drucker Diagnostics by Joshua Crouse, Regulatory Compliance Specialist

Quality management system registered to ISO 13485:2016

Form 1128 (Rev A)