Declaration of Conformity

Manufacturer: KD Scientific
Address: 84 October Hill Road, Holliston, Massachusetts 01746-1388, USA

We herewith declare that the following products:

Product Name: KDS Legato 100 and 950 Series Syringe Pumps
Models: KDS Legato 100 Series (78-81XX)
KDS Legato 950 Series (78-89XX)

Are in conformity of the following applicable European regulations and directives:

- 2014/35/EU Low Voltage directive (LVD)
- 2014/30/EU Electromagnetic Compatibility directive (EMC)
- 2012/19/EU Waste electrical and electronic equipment directive (WEEE)
- 2011/65/EU Restriction on the use of certain hazardous substances directive (RoHs)

Standards used to demonstrate conformity include:

- EN 61326-1:2013 Electrical equipment for measurement, control and laboratory use – EMC requirements
- EN 61010-1:2010 Safety requirements for electrical equipment for measurement, control and laboratory use, General Requirements
- EN 50581:2012 Assessment of products with respect to RoHS

1Classified as belonging to equipment categories 8 or 9
2Classified and tested as class A equipment in accordance with CISPR 11 definition in a basic electromagnetic environment. This equipment has also been tested and found to comply with the limits for a class A digital device, pursuant to CFR Title 47 part 15 of the FCC rules.

EMC and Safety compliance were evaluated by Intertek/ETL Semko

Reference test reports file numbers: 100602821 Box-001, -001a
100179785 Box –001, -003
100179785 Box –005a, -005b

I, the undersigned, hereby declare that the equipment specified is in conformity with the relevant harmonised Union legislation. Signed for and on behalf of Harvard Apparatus at Holliston, USA

Signed: Mark Davis
Director of Engineering

Date: October 09, 2017

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