



# CE Declaration of Conformity

**We:** Labconco Corporation  
8811 Prospect Avenue  
Kansas City, MO 64132 USA  
Ph. 816-333-8811 Fax. 816-363-0130  
labconco@labconco.com

***being the representative party solely responsible for this certification, hereby declare that the following products:***

Equipment: Laboratory Equipment – CentriVap Concentrators, Consoles & Systems

Type, Model: CentriVap DNA Concentrators 79700xx, where xx= 11, 30, 35, 37  
CentriVap Benchtop Concentrators 78100xx, where xx= 30, 31, 32, 34, 35, 36, 38, 39, 40  
CentriVap Acid Resistant Concentrators 78100xx, where xx= 17, 33, 37, 41  
CentriVap Refrigerated Concentrators 73100xx, where xx= 30, 31, 32, 34, 35, 36, 37, 38, 39, 40, 41, 42  
CentriVap Mobile Console Systems 78120xx, where xx= 11, 12, 14, 15, 35, 36, 37, 38  
CentriVap Concentrator Systems  
Aqueous Systems 79820xx, where xx= 11, 12, 35, 37  
Acid Systems 79830xx, where xx= 13, 14, 15, 35, 37  
Solvent Systems 7995800, where xx= 11, 12, 35, 37  
Proteomic Systems 79896xx, where xx= 01, 02, 35, 37  
CentriVap Complete 73150xx, where xx= 30, 31, 32, 33, 40, 41, 42, 43, 60, 61, 62, 63, 70, 71, 72, 73  
CentriVap Cold Trap 78110xx, 74600xx, or 73850xx, where xx= 20, 21, 30, 31, 35, 36, 37, 38, 40, 41

Manufacturer: Labconco Corporation  
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***are in accordance with the following directives:***

2014/30/EU	Electromagnetic Compatibility Directive	2015/863/EU	RoHS3 Directive
2014/35/EU	Low Voltage Directive		
2006/42/EU	Machinery Directive		
2011/65/EU	RoHS Directive		

***are in conformity with the standards listed below:***

EN61010-1:2010	EN61000-4-3:2020
EN61326-1:2013	EN61000-4-4:2012
EN55011:2016+A1:2017+A11:2020	EN61000-4-5:2014+A1:2017
EN61000-3-2:2014	EN61000-4-6:2014
EN61000-3-3:2013	EN61000-4-11:2020
EN61000-4-2:2009	EN63000:2018

***when installed and operated in accordance with the manufacturers installation and operating instructions.***

**Brandon Gray**  
Vice President of Engineering  
Labconco Corporation  
Kansas City, MO USA

29 JAN 2024  
Date of issue



# UKCA Declaration of Conformity

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***being the representative party, in accordance with UK government guidance, solely responsible for this certification, hereby declare that the following product:***

Equipment: Laboratory Equipment – CentriVap Concentrators, Consoles & Systems

Type, Model: CentriVap DNA Concentrators 79700xx, where xx= 11, 30, 35, 37  
CentriVap Benchtop Concentrators 78100xx, where xx= 30, 31, 32, 34, 35, 36, 38, 39, 40  
CentriVap Acid Resistant Concentrators 78100xx, where xx= 17, 33, 37, 41  
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Aqueous Systems 79820xx, where xx= 11, 12, 35, 37  
Acid Systems 79830xx, where xx= 13, 14, 15, 35, 37  
Solvent Systems 7995800, where xx= 11, 12, 35, 37  
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CentriVap Complete 73150xx, where xx= 30, 31, 32, 33, 40, 41, 42, 43, 60, 61, 62, 63, 70, 71, 72, 73  
CentriVap Cold Trap 78110xx, 74600xx, or 73850xx, where xx= 20, 21, 30, 31, 35, 36, 37, 38, 40, 41

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***the object of the declaration described above is in conformity with the relevant UK Statutory Instruments (and their amendments):***

S.I. 2016/1091 Electromagnetic Compatibility Directive  
S.I. 2016/1101 Low Voltage Directive  
S.I. 2008/1597 Machinery Directive  
S.I. 2012/3032 RoHS Directive

***are in conformity with the standards listed below:***

EN61010-1:2010	EN61000-4-3:2020
EN61326-1:2013	EN61000-4-4:2012
EN55011:2016+A1:2017+A11:2020	EN61000-4-5:2014+A1:2017
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