ATTESTATION OF CONFORMITY

Directive(s):	*Electromagnetic Compatibility Directive (2017/746)* *Electromagnetic Compatibility Regulations 2016*
Attestation No.:	*22070149*
Applicant / Holder:	*BLUE-RAY BIOTECH CORP.*
Address:	*5F., No. 2, Aly. 2, Siwei Ln., Zhongzheng Rd., Xindian Dist.,* *New Taipei City 23148, Taiwan (R.O.C.)*
Trade Name:	*BLUE-RAY BIOTECH*
Product / Test Item:	*EzDrop*
Model / Type Reference:	*EzDrop 1000, EzDrop 1000C*

The submitted sample(s) have been tested with the following standard(s) and found to be in compliance with the essential requirements of the Directive(s):

Standard(s)	
EN 55011:2016+A11:2020 (Class A)	EN IEC 61326-1:2021, BS EN IEC 61326-1:2021
BS EN 55011:2016+A11:2020 (Class A)	EN IEC 61326-2-6:2021, BS EN IEC 61326-2-6:2021
CISPR 11:2015+A1:2016+A2:2019	IEC 61000-4-2:2008, BS EN 61000-4-2:2009
EN IEC 61000-3-2:2019+A1:2021	IEC 61000-4-3:2020, BS EN IEC 61000-4-3:2020
BS EN IEC 61000-3-2:2019+A1:2021	IEC 61000-4-4:2012, BS EN 61000-4-4:2012
EN 61000-3-3:2013+A2:2021	IEC 61000-4-5:2014/AMD1:2017,
BS EN 61000-3-3:2013+A2:2021	BS EN 61000-4-5:2014+A1:2017
	IEC 61000-4-6:2013, BS EN 61000-4-6:2014
	IEC 61000-4-8:2009, BS EN 61000-4-8:2010
	IEC 61000-4-11:2020, BS EN IEC 61000-4-11:2020

The referred test report(s) show that the product fulfills the essential requirements set out in the Directive and the Regulation. On this basis, together with the manufacturer's own documented production control, the manufacturer or his European authorized representative can in his EC or UKCA Declaration of Conformity verify compliance with the Directive and the Regulation. The CE or UKCA marking could be affixed only when all the relevant and effective EC Directives or UK Regulations are complied with.

CE UKCA

Kero Kuo / EMC & RF Manager
2022-11-24

Cerpass Technology Corporation

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