ATTESTATION OF CONFORMITY

Attestation Number :	AOCRSC201012004-01
Date of Issue:	2020-11-02
Product:	Deep Tissue Massager
Model(s):	QL/MINI.L2-A
Multiple Model:	QL/MINI.L2-B, QL/MINI.L2-C, QL/MINI.L2-D, QL/MINI.L2-E, QL/MINI.L2-F, QL/MINI.L2-G, QL/MINI.L2-H, MGPC-002
Brand:	Beoka
Manufacturer & Address:	Sichuan Qianli-beoka Medical Technology Inc. Longtan industrial park, second section of eastthird ring road, chenghua district, chengdu，Sichuan, China

Bay Area Compliance Laboratories Corp. (ChengDu) hereby declares that the submitted sample(s) of the above equipment has been tested for CE-marking and in accordance with the following Standards:

Harmonized Standards	Test Report Number
EN60335-1:2012+A11:2014+A13:2017 EN60335-2-32:2003+A1:2008+A2:2015	RSC201012004-3

CE Mark is permitted only after all applicable requirements are met in accordance with the European Union Rules, including the manufacturer's issuance of a "Declaration of Conformity. The Declaration of Conformity is issued under sole responsibility of manufacturer. This attestation is specific to the standard(s) stated above and compliance with additional standards and/or European directives are applicable.

Attestation by: Alice Liu

Safety Director

Signature

COMPLIANCE TESTING AND VERIFICATION SERVICES BACL

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