CERTIFICATE

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Of Conformity
Low Voltage Directive 2014/35/EU

Registration No.: ATGZAHS190516003

Report No.: GZAHS190516003-01

Applicant	:	**Beijing LaserTell Medical Co., Ltd.** Block1, No.12 Jingsheng South 2nd Rd., JQ Science Park, Tongzhou District, Beijing, China	
Product	:	**Beauty Devices**	
Identification	**Model No.**	:	**AlexMED, AlexMED Pro, DepiMED, DepiMED Pro, NdMED, ClearMED, FemiMED, VenaMED, EpiMED, EpiMED Pro, TruMED, ExtrMED, iMED, PluShape, PluShape Pro, HydroCoolol, Faceage, 4Deepoo**
	Trade Mark	:	**LaserTell**
	Rating	:	**AC230V, 50/60Hz, 15A**
Test Standards	:	**EN 60335-1:2012+A11:2014+A13:2017** **EN 62233:2008**	

The certificate of conformity is based on an evaluation of a sample of the above mentioned product. Technical report and documentation are at the applicant's disposal. This is to certify that the tested sample is in conformity with Low Voltage Directive 2014/35/EU relating to electrical equipment designed for use within certain voltage limits. The certificate does not imply assessment of the series-production of the product. The applicant of the certificate is authorized to use this certificate in connection with EU declaration of conformity specified in Article 15 and Annex IV of the Directive.

Anbotek (GZ) Compliance Laboratory Limited
Anbotek Product Safety
* Approved *

Certified by

Terry Tian

Terry Tian
Manager

Jun. 11, 2019
Date

CE **The CE Marking may only be used if all relevant and effective EU Directives are complied with** CE

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