CERTIFICATE

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Of Conformity
EU Council Directive 2014/30/EU
Electromagnetic Compatibility

Registration No.: ATSZAHE190516005

Report No.: SZAHE190516005-01

Applicant	:	**Beijing LaserTell Medical Co., Ltd.**		
		Block1, No.12 Jingsheng South 2nd Rd., JQ Science Park, Tongzhou District, Beijing, China		
Product Identification	:	**Beauty Devices**		
		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 55014-1: 2017** **EN 61000-3-2: 2014** **EN 61000-3-3: 2013** **EN 55014-2: 2015**		

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 all provisions of Annex II of Council Directive 2014/30/EU, in its latest amended version, referred to EMC Directive. The certificate does not imply assessment of the production and does not permit the use of Lab's logo. The applicant of the certificate is authorized to use this certificate in connection with EU declaration of conformity to Article 15 of the Directive.

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

Certified by

Sally Zhang

Sally Zhang
Manager

Jun. 13, 2019
Date

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

Shenzhen Anbotek Compliance Laboratory Limited
1/F, Building D, Sogood Science and Technology Park, Sanwei community, Hangcheng Street, Bao'an District, Shenzhen, Guangdong, China.518102
Tel: (86)755-26066440 Fax: (86)755-26014772
Http://www.anbotek.com Email: service@anbotek.com