

## **Certification of Conformity**

## **EU-MEDICAL DEVICES DIRECTIVE-93/42/EEC**

Registration No.: ENC1908168GZ19

Applicant : Beijing LaserTell Medical Co., Ltd.

Applicant Address : Block1, No.12 Jingsheng South 2<sup>nd</sup> Rd., JQ Science Park,

Tongzhou District, Beijing, China.

Product Designation : Diode Laser & RF Systems

Model Number : AlexMED; AlexMED Pro; DepiMED; DepiMED Pro; 4Deepoo;

PluShape; PluShape Pro

Manufacturer : Beijing LaserTell Medical Co., Ltd.

Manufacturer Address : Block1, No.12 Jingsheng South 2<sup>nd</sup> Rd., JQ Science Park,

Tongzhou District, Beijing, China.

The submitted products have been tested by us with the listed standard and found in compliance with the Medical Devices Directive 93/42/EEC and following European Standards:

Directive	Applied Standards	Test Report No.
EMC Directive	EN 60601-1-2:2015	ENC1908168GZ19E1
	EN 61000-3-2:2014	
	EN 61000-3-3:2013	
Safety Directive	EN 60601-1:2006+A11:2011+A1:2013 +A12:2014	ENC1908168GZ19L1
	EN 60825-1:2014	

This certificate is based on an evaluation of a sample of the above mentioned product. Technical report and documentation are at the licence applicant's disposal. This certificate does not imply assessment of the series-production of the product. The CE markings as shown below can be affixed on the product after preparation of necessary technical documentation.

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Ray Zhou / General Manager Date of Issue: Aug. 23, 2019

## East Notice Certification Service Co., Ltd.

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