CERTIFICATION OF FDA REGISTRATION

This certifies that:

ZHEJIANG RECI LASER TECHNOLOGY CO., LTD.

Scientific Research Plant No. 2, Jintang North Road No. 2, Eastern New District, Wenling City, Taizhou City, Zhejiang Province, China

has been filed according to reporting requirements in Title 21, Code of Federal Regulations (CFR), Part 1002. Submission has been assigned an informal subject title below after "Purpose:". Submission has been assigned an ACCESSION NUMBER which can be used by you and FDA to identify submission

Accession Number : 2111483-000

Product name : CO2 laser tube

Purpose model(s) : W4, W1, T07, T4, W8, T08, T6, W2, T2, T1, W6.

This certificate affirms that the above stated facility is registered with the U.S. Food and Drug Administration pursuant to Title 21, Code of Federal Regulations (CFR), Part 1002, such submission having been verified as effective by OUA as of the date hereof, and OUA will confirm that such registration remains effective upon request and presentation of this certificate until the expiration of one year from the date hereof, unless terminated after issuance of this certificate. OUA makes no other presentations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. OUA assume no liability to any person or entity in connection with the foregoing. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. OUA is not affiliated with the U.S. Food and Drug Administration.
Please note that your firm is required to submit an Annual Report to CDRH every year by September

Judy Yi /Chief engineer
Issued Date: 09/08/2021
Expiration Date: 08/31/2022

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