FA6

Certification of Conformity

EU-MEDICAL DEVICES DIRECTIVE-93/42/EEC

Registration No.: ENC1911052GZ09

Applicant : Dermax Medical Technology (Hebei) Co., Ltd

Applicant Address : 2801, CAIKU International Commercial Port, no. 41, South Zhai

Ying Street, Yuhua district, Shijiazhuang City, Hebei Province

Product Designation : Micro Cannula

Model Number : 18G, 21G, 23G, 25G, 27G, 30G

Brand Name : Cozysculpt

Classification of Medical Devices: Class III

Manufacturer : Dermax Medical Technology (Hebei) Co., Ltd

Manufacturer Address : 2801, CAIKU International Commercial Port, no. 41, South Zhai

Ying Street, Yuhua district, Shijiazhuang City, Hebei Province

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:

Requirement	Applied Standards	Test Report No.
DELAYED-TYPE HYPERSENSIVITY	EN ISO 10993-1:2009	ENC1911052GZ09E1
	EN ISO 10993-10:2013	
	EN ISO 10993-12:2012	

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.

CE

Ray Zhou / General Manager Date of Issue: Nov. 5, 2019

East Notice Certification Service Co., Ltd.

1/F, Haohui Commercial Building, Zhuji Street, Dongpu Town, Tianhe District, Guangzhou City
Tel:+86-020-8256 8534 Fax:+86-020-8256 8534 E-mail:enc@enc-lab.com Web: www.enc-lab.com