FDA

Fiscal Year 2021

CERTIFICATION OF REGISTRATION

This certifies that:

Name: DE COREMATRIX CO.LTD.

Add: No.40 NanBeigang Road, HuKou County,Jiujiang, Jiangxi, 332599, CHINA

has completed the FDA Establishment Registration (as manufacturer and foreign exporter) and Device Listing with the US Food & Drug Administration, through

Registration/FEI Number:

Device Listing #:10080077

Listing No	Code	Premarket Submission	Device Name
D432913	EIH	K192262	POWDER, PORCELAIN

ABmed will confirm that such registration remains effective upon request and presentation of this certificate until the end of the year stated above, unless said registration is terminated after issuance of this certificate. ABmed makes no other representations 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. This certificate does not denote endorsement or approval of the certificate - holder's device or establishment by the U.S Food and Drug Administration.

ABmed assumes no liability to any person or entity in connection with foregoing.

Date of verification: Jan.12, 2021

Date of expiration: Dec. 31, 2021

SH OFFICE

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