

SHT-LAB



CERTIFICATION OF REGISTRATION

2020

SHT2003044FD

This certifies that:

GUIZHOU BOCAI MEDICAL DEVICES CO., LTD.

**Honghuagang district medical and health industrial park, ZUNYI, Guizhou,
563099, CHINA**

Was registered with US Food and Drug Administration, Center for Devices and Radiological Health, pursuant to the Code of Federal Regulations 21 CFR 807, by Shenzhen SHT Testing Technology Co., Ltd.

Owner / Operator Number:	10065023
Device Listing #:	See Annex
Expiration Date:	Dec. 31, 2020

SHT will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate.

SHT 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.

Pursuant to 21 CFR 807.39, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding." 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, SHT is not affiliated with the U.S. Food and Drug Administration.



Mar. 29, 2020

(Date)

Ryan MA
Lab Supervisor


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Annex to Device Listing #:

Proprietary Name	Product Codes	Device Class	Listing Number	Establishment Operations
Disposable medical masks	KHA	1	D380538	Manufacturer



Mar 29, 2020
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(Date)
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Ryan MA
Lab Supervisor