



The 3rd Party Certificate of  
FDA Medical Device Registration

**Note:**

This file is Not being issued by FDA. We, SFT, as the 3rd party, produce it, intended to facilitate customer display & transmit information. The following contents, FDA registered Facility/Owner/Operator&FDA listing Medical Device, are excerpted from database at [www.fda.gov](http://www.fda.gov).

**Establishment:**

Qingdao Kutesmart Co.,Ltd

No.17 Redcollar Street, District Jimo, Qingdao, Shandong,China 266200

Registration Number / FEI Number\*:

\* Firm Establishment Identifier (FEI) should be used for identification of entities within the imports message set

Status: **Active**

Date of Registration Status: **2020**

**Owner/Operator**

Qingdao Kutesmart Co.,Ltd

No.17 Redcollar Street, District Jimo, Qingdao, Shandong,China 266200

Owner/Operator Number: 10063542

**Official Correspondent**

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**U.S. Agent**

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**Devices Listing Information**

Proprietary Name	Product Codes	Device Class	Listing Number	Establishment Operations
disposable medical mask	LYU	1	D37****	Manufacturer
Disposable Face Mask	LYU	1	D37****	Manufacturer
Protective Clothing	OEA	1	D37****	Manufacturer
Insulating Clothing	OEA	1	D37****	Manufacturer

⚠ Please careful protect your Listing Number.

Professional FDA Registration Services, by Shanghai Shifu Testing Service Co., Ltd.

More details on the website: <http://www.sft-lab.com>.

Need help? Contact us, SFT, at +86(021) 51300821&[sales@sft-lab.com.cn](mailto:sales@sft-lab.com.cn)

FDA CERTIFICATE NUM: **SFT20MAR124C**