



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:**

[JIANGSU YIMAO FILTER MEDIA CO.,LTD](#)

North side of Fudan Rd,Sucheng Economic and Development Zone Suqian  
Jiangsu, CN 223800

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**

[JIANGSU YIMAO FILTER MEDIA CO.,LTD](#)

North side of Fudan Rd,Sucheng Economic and Development Zone Suqian  
Jiangsu, CN 223800

Owner/Operator Number: [10062630](#)

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**Devices Listing Information**

Proprietary Name	Product Codes	Device Class	Listing Number	Establishment Operations
Face Mask	LYU	1	D372175	Manufacturer

⚠ Please careful protect your Listing Number.

Approved by: Reilly

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](mailto:sft-lab.com).cn

FDA CERTIFICATE NUM: [SFT20FEB006C](#)

