Certificate of Registration

This certificate of Registration certifies that:

SHANGHAI OGO MEDICAL INSTRUMENT CO., LTD.

5F BUILDING 1, NO.2 LANE 1515, NORTH HUIFENG ROAD, FENGXIAN DISTRICT, SHANGHAI, 201499, CHINA

has registered with the US Food and Drug Administration pursuant to Title 21 of the United States Code of Federal Regulations.

Owner Operator Number: 10071208

Listed Device

MSH D394867 Respirator, Surgical

US Agent: Willow Glen Consultancy LLC
Willow Glen Number: WG2071055
Expiration Date: December 31, 2020

This certificate affirms that the above-named facility is registered with the US FDA pursuant to the regulations required by the US laws. This registration has been verified as effective by Willow Glen Consultancy as of the date below, unless such registration has been terminated after issuance of this Certificate. Willow Glen Consultancy makes no additional representations or warranties, nor does this certificate carry any to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Willow Glen Consultancy assumes no liability to any person or entity in connection with the foregoing, nor does the U.S. FDA recognize a certificate of registration issued by Willow Glen Consultancy.

Willow Glen Consultancy is a private agent not affiliated with the U.S. Food and Drug Administration.

Amanda Ou, Operation Director
Willow Glen Consultancy LLC
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Date: April 20, 2020

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