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Device Classification Name	Mask, Surgical ²²
510(K) Number	K173062
Device Name	Non Woven Face Mask (Models: VQN0185W (Earloop) And VQN0185B (Ties))
Applicant	V&Q Manufacturing Corporation #B1614 Optical Valley Time Square, Wuhan, CN 430074
Applicant Contact	Jacky Jin
Correspondent	Feiying Drug & Medical Consulting Technical Service Group B-3F 3005, Bldg.1, Southward Ruifeng Business Center, No 22 Guimiao Road Shenzhen, CN 518000
Correspondent Contact	Tracy Che
Regulation Number	878.4040 ²³
Classification Product Code	FXX ²⁴
Date Received	09/28/2017
Decision Date	01/25/2018
Decision	Substantially Equivalent (SESE)
Regulation Medical Specialty	General & Plastic Surgery
510k Review Panel	General Hospital
Summary	Summary ²⁵
Type	Traditional
Reviewed By Third Party	No
Combination Product	No

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Page Last Updated: 07/06/2020

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