Notification of New Device Establishment Registration

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Center for Devices and Radiological Health

10903 New Hampshire Ave., WO66 Room 2621

Silver Spring, Maryland 20993-0002

July 2, 2020

Name of Official Correspondent	Jerry Yang
Address of Official Correspondent	Room 201, 1st building No.5 Industrial avenue, Shipai Town
	Dongguan, Guangdong, 523330
	CHINA
Registration Number	3016676280
Owner Operator Number	10064273
Dear Sir or Madam,	
We have received your registration and listing information for the following medical device establishment	
Establishment Name	DONGGUAN FORES BIOTECHNOLOGY CO., LTD
Establishment Address	Room 201, 1st building No.5 Industrial avenue, Shipai Town
	Dongguan, Guangdong, 523330
	CHINA

The information submitted has been processed and entered into the FDA Registration and Device Listing Database. Your device establishment is now considered registered. You will be notified of your official registration number within 90 days.

Once you receive a registration number, you are required to re-register on an annual basis from October through December. Failure to re-register every year will invalidate your registration and result in your device establishment and listing information being removed from the FDA Medical Device Registration and Listing Web site.

ATTACH

Device Listing		
Device#	**Code**	**Device Name**
D388241	QKR	Face mask (except N95 respirator) for general public/healthcare personnel per IIE guidance (INVISISMART FACE MASK)