FDA Registration Information

Proprietary Name:	MDU/Rx
Classification Name:	PURIFIER, AIR, ULTRAVIOLET, MEDICAL
Product Code:	FRA
Device Class:	2
Regulation Number:	880.6500
Medical Specialty:	General Hospital
Registered Establishment Name:	HGI INDUSTRIES INC
Registered Establishment Number:	3008156690
Premarket Submission Number:	K133800
Owner/Operator:	HGI Industries Inc
Owner/Operator Number:	10070115
Establishment Operations:	Manufacturer

https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRL/rl.cfm?lid=668968&lpcd=FRA