PROTECH LEADED EYEWEAR INC,
DBA PROTECH MEDICAL

1360 North Killian Drive, Unit 2
Lake Park, FL 33403

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FDA REGISTRATION

1. GENERAL

ESTABLISHMENT

Protech Leaded Eyewear, Inc.

Business Trade Name:
Protech Medical
1360 North Killian Drive, STE 2
Lake Park, FL 33403

Registration Number: 1063300
FEI Number*: 1000488948
Status: Active
Date of Registration Status: 2022

Owner/Operator Number: 10025412
Owner/Operator:
Protech Leaded Eyewear, Inc.
1360 North Killian Drive, STE 2
Lake Park, FL 33403

2. PRODUCT LISTINGS

APRONS

Classification Name: Apron, Protective
Product Code: IWO
Device Class: 1
Regulation Number: 892.6500
Medical Specialty: Radiology

BARRIER

Classification Name: Barrier, Control Panel, X-Ray, Movable
Product Code: IWX
Device Class: 1
Regulation Number: 892.6500
Medical Specialty: Radiology

EYEWEAR

Classification Name: Shield, Eye, Radiological
Product Code: IWS
Device Class: 1
Regulation Number: 892.6500
Medical Specialty: Radiology

GLOVES

Classification Name: Radiographic Protection Glove
Product Code: IWP
Device Class: 1
Regulation Number: 892.6500
Medical Specialty: Radiology

NITRILE GLOVES

Classification Name: Polymer Patient Examination Glove
Product Code: LZA
Device Class: 1
Regulation Number: 880.6250
Medical Specialty: General Hospital

Kindly note that all information contained herein is publicly available information which can be found on the FDA's website:
* Access: https://www.fda.gov/medical-devices/device-registration-and-listing/search-registration-and-listing
* Select: "Search the Registration & Listing Database"
* Input: "Protech Leaded Eyewear" into the Establishment or Tradename Section and click on the Search Button.
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