



PROTECH LEADED EYEWEAR INC,  
DBA PROTECH MEDICAL

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

Email(USA): sales@protechmed.com  
Email(Intl): global@protechmed.com  
Email (LatAm): LatAm@protechmed.com  
Phone: 561.627.9769  
Web: www.protechmed.com

## 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:** 2025

**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  
**Registration Type:** IWO  
**Product Code:** Manufacturer, Contract manufacturer  
**Device Class:** 1  
**Regulation Number:** 892.6500  
**Medical Specialty:** Radiology

#### BARRIER

**Classification Name:** Barrier, Control Panel, X-Ray, Movable  
**Registration Type:** Manufacturer, Contract manufacturer  
**Product Code:** IWX  
**Device Class:** 1  
**Regulation Number:** 892.6500  
**Medical Specialty:** Radiology

#### GLOVES

**Classification Name:** Radiographic Protection Glove  
**Registration Type:** Specification Developer  
**Product Code:** IWP  
**Device Class:** 1  
**Regulation Number:** 892.6500  
**Medical Specialty:** Radiology

#### EYEWEAR

**Classification Name:** Shield, Eye, Radiological  
**Registration Type:** Manufacturer, Contract manufacturer  
**Product Code:** IWS  
**Device Class:** 1  
**Regulation Number:** 892.6500  
**Medical Specialty:** Radiology

#### SCATTERGUARD DRAPES

**Classification Name:** Drape, Surgical, Exempt  
**Registration Type:** Specification Developer  
**Product Code:** PUI  
**Device Class:** 2  
**Regulation Number:** 878.4370  
**Medical Specialty:** General and Plastic Surgery

#### NITRILE GLOVES

**Classification Name:** Polymer Patient Examination Glove  
**Registration Type:** Specification Developer  
**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.

NOTICE: The FDA does not market nor endorse company marketing or claims. Be cautious about those including the FDA logo on their literature.