

PROTECH LEADED EYEWEAR INC. DBA PROTECH MEDICAL

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EU EYEWEAR DECLARATION OF CONFORMITY

This declaration of conformity is issued under the sole responsibility of Protech Leaded Eyewear, Inc. DBA Protech Medical. Declares that the Radiation Protective Eyewear made with 0.75mm or 0.50mm Pb lenses, "PPE" described hereafter:

C/T **LENS TYPE** LENS CHARACTERISTICS **KV RANGE** MM PB 0.50 / 0.75mm Pb 65mm 400B, 73mm 600B, 73mm 800B, 80mm 400B 2.5mm+ >= 150kV Plano 0.50 / 0.75mm Pb Semi-Finished Rx 200B, 400B, 600B, 800B 2.5mm+ >= 150kV 4B - Main Lens; 6B - Main Lens; 8B - Main Lens 2.5mm+ 0.50mm Pb **Progressive** >= 150kV **Side Shields** Vinyl Side Shields N/A >= 150kV 0.50mm Pb **Face Shields** Acrylic Panel 0.12mm+ Pb N/A >=120kV

EN 61331-1:2014 AND EN 61331-3:2014

Glasses or Goggles (heavy duty)

Protech Brand: RE-90; RE-42; RE-FGUARDA; RE-FGUARDB; RE-705; RE-70E; RE-COMET; RE-COM; RE-OG220; RE-XGUARD; RE-53; RE-53E; RE-MALIB; RE-RETRO; RE-002; RE-003; RE-007; RE-008; RE-553; RE-553S; RE-557S; RE-662S; RE-98; RE-99; RE-99R; RE-99SM; RE-994I; RE-994I53; RE-994IA; RE-BABY; RE-AIRB; RE-CENT; RE-CIRC; RE-ONYX; RE-MAKO; RE-RAZER; RE-TBABY; RE-VAL; RE-EDGE; RE-RYK; RE-ATLAS; RE-ASPEN; RE-LOTUS; RE-LEG; RE-JUN; RE-LOND; RE-ATH; RE-LAT

Abaco Brand: AB-DOCK: AB-JES: AB-KAI: AB-TIKI: AB-WAI

Costa Brand: CO-ALE; CO-ARA; CO-BFIN; CO-BRD330; CO-BRD320; CO-FER; CO-OCR4; CO-SEA; CO-SUL; CO-TALY; CO-TYB

Liberty Brand: LBT-MX30S; LBT-MX30; LBT-MX31S; LBT-BIKER

Nike Brand: N-7116; N-7117; N-7118; N-7245; N-7284; N-7286; N-ADR; N-BR; N-BRF; N-BRS; N-MAV; N-PRMR; N-RABID; N-WAVE

Oakley Brand: OA-CROSSZ; OA-FIVESQ; OA-HOL; OA-JUP; OA-LITE; OA-STRGHT; OA-TRILL

Ray Ban Brand: RB-2016; RB-3534; RB-4105; RB-4165; RB-4181; RB-7047 Wiley-X Brand: WX-AIRR; WX-BRICK; WX-BOSS; WX-CON; WX-P17

Gucci Brand: GG-0372; GG-0752; GG-0758

Face Masks

450FPM; 300GOG; 400PS; 350FLP

Leaded Eyewear ("PPE") described above is in conformity with the provisions of Regulation (EU) 2016/425 and the models satisfy the requirements of the manufacturer's technical and quality management specifications with testing based upon IEC 61331-1:2014 and IEC 61331-3:2014. This is identical to the PPE which Shirley Technologies (Europe) Limited, Notified Body 2895 (Port Tunnel Business Park, Office 13 Unit 21, Dublin 17, ROI), performed the EU Module B type-examination on and issued the EU type-examination Certificate SH00792. The PPE is subject to the conformity assessment procedure set out in Regulation (EU) 2016/425, Module D (Certificate #SH00654) under surveillance of the notified body Shirley Technologies (Europe), Limited, notified body number 2895, performed at Protech Leaded Eyewear, Inc. DBA Protech Medical, 7960 Central Industrial Drive, Suite 125, West Palm Beach, FL 33404 USA.

Signed for and on behalf of: Protech Leaded Eyewear, Inc. DBA Protech Medical, at 7960 Central Industrial Drive, Suite 125, West Palm Beach, FL 33404 USA.

Jarrod Parasmo, President Signature Printed Name Date

July 7th, 2025