

User's Manual of Safety Goggle

Safety goggle is non-sterile disposable anti-fog goggle.

- 1. Brand: Refine
- 2. Product name: Safety Goggle 3. Material: PVC 70% PC 30%
- 4. Model: EP01
- 5. Composition: composed of protective cover and fixing device made of polymer material. Non-sterile, single use.
- 6. Applied standards: GB 14866-2006 / EN166:2001 / ANSI/ISEA Z87.1-2015 AS/NZS 1337.3:2010 / CSA Z94.3-2015
- 7. Intended use: Safety Goggle is common eye protection equipment, which are used for protecting against droplets and splashes of liquids. Recommended industry: Facility Sanitation, Food Processing, Food Safety.
- 8. Product instruction: The product and the protective packaging should be checked before using. Stop using it if there is any damage.
- 9. Precautions: 9.1 One-time use only.
- 9.2 Use in accordance with the instructions.
- 9.3 Never use it when damages are found.
- 9.4 Stay away from chemicals.
- 9.5 The eye protector is not intended to protect against high speed particles.
- 10. Storage: Please store in clean, dry and ventilated indoor place, with relative humidity (10%-93%), Atmospheric pressure for storage (70kPa-106kPa), temperature (-20°C- +40°C), avoid corrosive gas.

Marning:

- a. Materials which may come into contact with the wearer's skin could cause allergic reactions to susceptible individuals:
- b. Scratched or damaged oculars should be replaced;
- c. Eye-protectors against high speed particles worn over standard ophthalmic spectacles may transmit impacts, thus creating a hazard to the wearer.
- d. These protectors are intended for indoor use where no optical radiation hazards exist.

Symbol instruction

| Symbol 符号 | Instruction 说明 | | |
|---------------|------------------------------------------------------------------|------------------------|--|
| <u></u> | Warning, Caution and Important ! Check the Instruction Manual | 注意! 查阅随机文件 | |
| <u>~</u> | Date of manufacture | 生产日期 | |
| 6 | Recovery | 可回收标志 | |
| | Keep dry | 防潮标识 | |
| 70 KPa 106 | Atmospheric pressure for storage | 存贮条件,大气范围为70kPa~106kPa | |
| -20°C - +40°C | Temperature limitation for storage | 存贮条件,温度范围为-20℃~+40℃ | |

| Symbol 符号 | Instruction 说明 | | | |
|-------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------|--|
| LOT | Lot number | 批次号 | | |
| Z87 D3 | According to ANSI/ISEA Z87.1-2015: D3-Special use, used to prevent Droplet / splash | 根据美国ANSI/ISEA Z87.1-2015标准的分类: D3-特殊用途,用于防护液滴/飞溅 | | |
| EC REP | EU Representative | 欧盟授权代表 | | |
| ^^ | Manufacturer | 制造商 | | |
| | Handle with care | 易碎产品,小心轻放 | | |
| i | Consult the accompanying documents | 查阅随机文件 | | |
| 2 | Single-Use | 一次性使用 | | |
| 10% | Humidity limitation for storage | 存贮条件,湿度范围为10%~93% | | |
| C€ | CE certification | 欧盟CE认证 | | |
| REFINE 1SN EN166 3 S | According to EN 166:2001, REFINE - manufacture LOGO 1 - Optical class (Class I) S - Symbol for increased robustness N - Symbol for Resistance to fogging of oculars EN 166 - Product Standards 3 - Field of use (Liquid Droplet/splash) S - Symbol for Increased robustness | | 根据欧盟EN 166:2001的防护分类: REFINE - 制造商商标 1 - 光学等级分类I类 S - 提高坚固性的符号(镜片) N - 防雾镜标识 EN 166 - 适用欧盟标准 3 - 防护液滴/飞溅 S - 提高坚固性的符号(镜架) | |
| ERC | EAC certification | | 海关联盟(俄罗斯)认证 | |
| MDEL | MDEL certification | | 加拿大MDEL认证 | |

医用护目镜使用说明书

医用护目镜是非无菌一次性使用的眼镜。

1. 产品品牌: Refine

2. 产品名称: 医用护目镜

3. 产品成分: PVC 70% PC 30%

4. 产品型号: EP01

5. 产品性能结构及组成:由高分子材料制成的防护罩和固定装置组成。非无菌提供,一次性使用。

6. 产品标准: GB 14866-2006 / EN166:2001 / ANSI/ISEA Z87.1-2015

AS/NZS 1337.3:2010 / CSA Z94.3-2015

7. 适用范围:用于医疗机构中检查治疗时起防护作用,阻隔体液、血液飞溅或泼溅。

产品禁忌症: 暂不明确。

- 8. 产品使用说明:本器械仅适用于其对应的适用范围,不应用于其他用途,用于其他用途将会损坏器械或者效果不好。应由专业人士进行操作使用,小心使用此器械。每次使用前,应对保护性包装和产品进行检查,如发现有破损的情况,该产品不应再继续使用。
- 9. 注意事项
- 9.1 本器械为一次性使用产品,请勿重复使用;
- 9.2 器械使用时请严格遵守使用说明且在使用过程中不得经受冲击、摔打;
- 9.3 如发现器械有破损现象,请勿使用;
- 9.4 器械应避免与化学药品并放;
- 9.5 本产品不防高速颗粒冲击。
- 10.贮存: 应贮存于清洁、干燥、通风,相对湿度为 10% ~ 93%,

大气压力为 $70kPa \sim 106kPa$,温度为 $-20^{\circ}C \sim +40^{\circ}C$ 的无腐蚀气体的室内环境中。

▲ 警告

- a. 与穿用者皮肤接触的材料可能对易过敏者造成过敏反应;
- b. 产品有刮伤或损坏的, 应予以更换;
- c. 佩戴了普通标准眼镜后,再使用该产品,用于高速颗粒的防护时,可能会通过普通眼镜传递冲击力,从而对佩戴者造成危害。
- d. 本产品适用于不存在光辐射危害的室内使用。



Guilin Refine Medical Instrument Co., Ltd.

No.8-3, Information Industrial Park, High-Tech Zone, Qixing District, Guilin, Guangxi, 541004, P.R. China

EC REP

REP MedPath GmbH

Mies-van-der-Rohe-Strasse 8, 80807 Munich, Germany

Shelf life: 2 Years

桂林市锐锋医疗 器械有限公司

产品名称: 医用护目镜

生产备案凭证编号: 桂桂食药监械生产备20170008号备案凭证 / 产品技术要求编号: 桂桂械备20200014号

备案人 / 生产企业/售后服务单位: 桂林市锐锋医疗器械有限公司

住所: 桂林市七星区高新区信息产业园8-3号

生产地址: 桂林市七星区高新区信息产业园8-3号;

桂林市七星区高新区信息产业园D-08号地块1#楼一层及2#楼

销售/售后服务电话/传真: 0773-7796686 邮箱: sales1@refine-med.com

网址: http://www.refine-med.com 使用年限: 2年

RF-EP1-M001 Edition: V1.6 Modify: 20200620

EU DECLARATION OF CONFORMITY

1. PPE (product, type, batch or serial number):

Safety Goggle, EP01, Category II, AZ611

2. Name and address of the manufacturer and, where applicable, his authorized representative:

Manufacturer: Guilin Refine Medical Instrument Co.,LTD.

Address of the manufacturer: No.8-3, Information Industrial Park, High-Tech Zone, Qixing District, Guilin, Guangxi, 541004, P.R.China

Authorised representative: MedPath GmbH

Address of the Authorised representative: Mies-van-der-Rohe-Strasse 8, 80807 Munich,
Germany

3. This declaration of conformity is issued under the sole responsibility of the manufacturer:

Manufacturer: Guilin Refine Medical Instrument Co.,LTD.

No.8-3, Information Industrial Park, High-Tech Zone, Qixing District, Guilin, Guangxi, 541004, P.R.China

4. Object of the declaration (Identification of PPE allowing traceability; where necessary for the identification of the PPE a colour image of sufficient clarity may be included):



Model: FP0

Brand: Refine Model: EP01 Lot No.: **AZ611** Production Date: June 2020

Safety Goggle

EP01:

Traceability Labeling:

- The object of the declaration described in point 4 is in conformity with the relevant Union harmonization legislation: Regulation (EU) 2016/425
- 6. References to the relevant harmonized standards used, including the date of the standard, or references to the other technical specifications; including the date of the specification, in relation to which conformity is declared:

Harmonised Performance Standard No(s): EN 166:2001

Technical specification No(s): RF-EP1-T001

Test Reports: C80132072R001

7. Where applicable, the notified body UL International (Netherlands) B.V. (European Notified Body No. 2821) performed the EU type-examination Module B and issued the EU type-examination certificate: 2821-PPE-0002.

Signed for and on behalf of Guilin Refine Medical Instrument Co.,LTD.

(place and date of issue): Guilin, 2020-06-07

Name: Jordan Chen, Title: Management representative (signature)

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