

TECHNICAL CONSTRUCTION FILE (FOPL/TCF/01)	Issue No.: 03 Dated :02-05-2022 Rev.No.:03 Rev Dt -24-09-2026	 Freedom Ophthalmic Pvt. Ltd. Plot No. 31, Phase -I, Sipcot Industrial Complex, Hosur – 635 126, Tamilnadu, INDIA
INSTRUCTIONS FOR USE (Doc No FOPL/IFU/01)	Section No.: 2.1 Page 1 of 7	

PMMA POSTERIOR CHAMBER IOLS:

ENGLISH

INSTRUCTIONS FOR USE APPLICABLE FOR THE FOLLOWING MODELS:

Trade / Brand Name	Generic Name	Models	Variants
FREEDOM LENS	PMMA Posterior Chamber Single Piece Intraocular Lens	PCC 503, PMC 524, PMC 553, PCC 603H, PMC 602, PMC 602H, PMC 652H, PSF 651	IOLs, APHAKIC, MONOFOC AL, ASPHERIC - OTHER
CLAR IOL		JMP 5025, JMP 5220, JMP 5525, JMP 6025H, JMP 6030, JMP 6030H, JMP 6530H, JMP 6535	
FREEDOM POWER	PMMA Posterior Chamber Single Piece Square Edge Aspheric Intraocular Lens	PMC 553SQA, PMC 603HSQA	
CLAR IOL		JMP 5525SQA, JMP 6025HSQA	

INTENDED USE:

It is an implant designed to be placed into the Posterior chamber for cataract surgery, suitable for replacing the crystalline lens in the correction of aphakic during surgical procedures in which the crystalline lens is removed.

INDICATIONS:

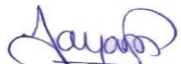
The Aphakic IOLs is indicated for patients to all age groups, who need a crystalline lens replacement after cataract extraction.

NOTES:

- This device should only be used by or under supervision of physicians trained for this process.
- This device is designed for single use only. Re-processing, re-sterilization and/or re-use may lead to failure of this device, and/or transmission of disease contracted from previous usage.
- Do not use this device:
 - For any purpose other than stated intended use.
 - If package is opened or damaged when received.
 - If date of use has expired.

CONTRAINDICATIONS/ RESIDUAL RISK:

The following are relative circumstances where the physician should consider whether implanting an intraocular lens does not create undue risk. Surgeons should explore the use of alternative methods of Aphakic correction and consider lens implantations only if alternative are deemed unsatisfactory to meet the needs of the patient.

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- Chronic severe Uveitis
- Rubella Cataract
- Concomitant severe Eye Disease
- Choroidal Hemorrhage
- IOL Dislocation
- Pseudoexfoliation Syndrome
- Retinal Pigment change
- In Cataract present in all age groups
- Posterior Capsule Opacification (PCO)
- Epithelial Dystrophy
- Massive vitreous loss
- Glaucoma problem
- Microphthalmos, Anirida
- Uveitis
- Myopia
- Eye trauma and connective tissue diseases
- Congenital cataract

POTENTIAL COMPLICATIONS:

The known complication may occur at the time of surgery or be the result of an on going post-operative process, mechanical and inflammatory may produce corneal de-compensation, Cystoid Macular Edema & Hyphema, Uveitis and Glaucoma may cause reduced vision and, in some cases, chronic pain in eye.

PRECAUTIONS:

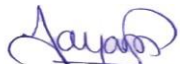
- Do not store the lens indirect sunlight & Keep away from freezing.
- Do not use if sterile pouch is opened & damaged.
- Only skilled Surgeons with experience in either viewing and or assisting numerous surgical implantations and successfully completed at least a course on IOL implantation should attempt implantation of these lenses.
- Pouch should be opened only under sterile conditions.
- Do not soak or rinse lens in solutions other than sterile balanced salt solution, sterile normal saline solute or equivalent of such.
- Do not attempt to re-sterilize this lens. Re-sterilization on this product has not been validated.
- Do not autoclave this lens.
- Handle the lens carefully, Locking forceps or needle holders should never be used to pick up lenses.
- Do not reshape the supporting structures. (Haptics).
- The device must be used by medical professional only.

ADVERSE EVENT:

The improper use / mis-use of lens may occur adverse event to patient eg Cystoid Macular Edema & Hyphema, endophthalmitis, anterior lens tissue on growth, lens dislocation, hypopyon, corneal edema, iritis, secondary glaucoma, secondary surgical intervention and glare\halos.

WARNINGS AND RESIDUAL RISKS:

- The effectiveness of UV-absorbing intraocular lenses in reducing the incidence of retinal disorders has not been established.
- The safety of the use of the Neodymium-YAG laser on IOLs with UV absorbing materials has not been established, the physicians is urged to use extreme caution in such cases where a patient with UV absorbing IOLs is treated with a Neodymium- YAG laser

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- The compression force exerted on the eye tissue by the lens is not established. The physician should have knowledge in selecting type of the lens depending on the eye dimensions.
- In the bag lenses should be used only when the posterior capsule is in good condition.
- Care should be taken to avoid breakage of haptic while injecting and inserting lens through the Scleral tunnel or small incision.
- **Don't re-sterilize the device.** Re-sterilization of the device will have a detrimental effect on its known properties thereby rendering it inappropriate for the intended use. Re-sterilization of Lens can also cause to change in mechanical & chemical properties.
- **Don't reuse the device.** if reuse of device can work as carrier for communicable disease, HIV, Chronic severe uveitis, Epithelial Dystrophy, Anirida, Hepatitis, contagious diseases, undue diseases to patient and/or user, Glaucoma problem, Rubella Cataract.
- Re-use of lens may not meet the intended use as lens may be blunt, Surgical difficulties at time of cataract extraction, which may increase the potential for complications (like persistent bleeding, significant iris damage, uncontrolled positive pressure or significant vitreous prolaps or loss)

MODE OF ACTION:

An IOL implant is an acrylic replacement of natural crystalline lens and take over the image focusing function of the eye. The IOLs are like a natural lens focus lights inside the eye. The devices can correct a wider range of vision problem than any other vision correction surgery. The IOLs create a single elongated focal point rather than several foci to enhance the depth of focus.

PREPARATION FOR DECONTAMINATION:

No particular requirement

APPLICATION OF THE BODY:

The device is implanted in eye primarily in capsular bag for visual correction. Thus, ocular tissue is coming in contact with device.

CLEANING AUTOMATED:

No cleaning is required before used of the device.

CLEANING MANUAL:

No cleaning is required before used of device.

DISINFECTION:

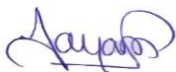
No disinfection required

MAINTENANCE:

No particular requirement

INSPECTION & FUNCTION TESTING:

Check for smooth functioning of medical device. Visually inspect for damage.

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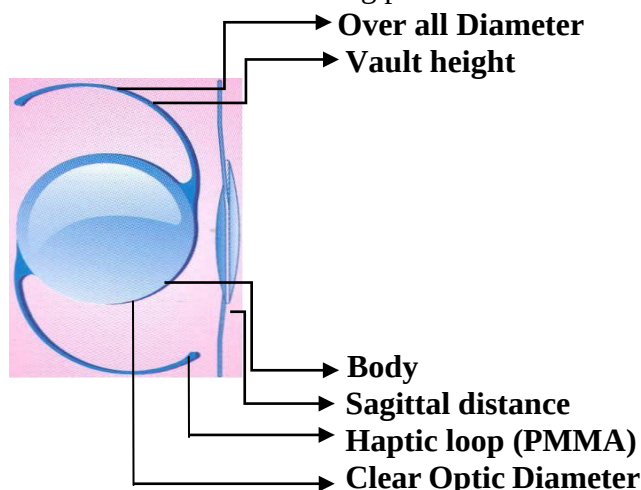
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ADDITIONAL INFORMATION:

Read instructions carefully before use.

DESCRIPTION OF COMPONENT PARTS:

The device consists of the following parts:



SYSTEM PREPARATION:

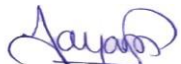
1. Check the label for manufactory and expiry dates (do not use the medical device after expiry)
2. Check for UDI number
3. Make sure the package is not damaged.
4. Visually examine the IOL to see if there are any damages on the loops.

INSTRUCTIONS FOR USE:

1. In a sterile environment, peel apart to open the pouch and remove the lens case.
2. Unscrew the lens case with care and grasp the lens optic using smooth edged forceps and lift.
3. Soak or rinse the lens in sterile balanced salt solution or their equivalent solution.
4. Ensure that the lens is in good conditions for optic and haptics surfaces for adherences of any particles.
5. Prior to implantation examine the lens for power, model, proper configuration, and optical surfaces.
6. During insertion carefully handle lenses by haptics portion only. To avoid breakage do not attempt to expand the haptics, flex the haptics out of the plane of the lens or twist or torque the lens.
7. Packing box contains extra labels. These are for convenience in maintaining and reporting records of implantable lenses during clinical investigation. One of the labels could be affixed in patients case sheet for future reference.

PATIENT INFORMATION:

The Surgeon performing the intervention must inform the patient about the complete procedure and all known complications and risks. The patient should be instructed to properly inform the doctor in charge about any side-effects after implantation.

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STORAGE:

Store in a cool, dry place, away from extreme temperatures. Expiry date showed on the pack indicates the period within which the device must be used. Store between 5°C to 40°C temperature & 35% to 65%RH away from moisture and direct heat.

RETURN OF DAMAGED PRODUCT:

Return the products in its original box identified by the LOT number, your purchase reference and reason for the return. The action shall be initiated to handle the product record as per the law of the land (IMDR 2017 under Drug and Cosmetic act 1945) and in compliance of the regulatory requirement of the destination of the product. The product recall procedure will be followed for the handling of this situation.

DISPOSAL OF DISCARDED PRODUCT AND PACKAGING:

Ensure safe and proper disposal of used/discarded product and packaging to avoid adverse effect to environment, children, and stray animals. The Disposal should comply with institutional guidelines for bio-hazardous medical waste prevailing in the country of use.

ORDERING INFORMATION:

Trade / Brand Name	Generic Name	Models	Diopter range
FREEDOM LENS	PMMA Posterior Chamber Single Piece Intraocular Lens	PCC 503, PMC 524, PMC 553, PCC 603H, PMC 602, PMC 602H, PMC 652H, PSF 651	-15 D to +40 D
CLAR IOL		JMP 5025, JMP 5220, JMP 5525, JMP 6025H, JMP 6030, JMP 6030H, JMP 6530H, JMP 6535	
FREEDOM POWER	PMMA Posterior Chamber Single Piece Square Edge Aspheric Intraocular Lens	PMC 553SQA, PMC 603HSQA	
CLAR IOL		JMP 5525SQA, JMP 6025HSQA	

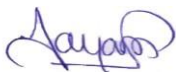
PACKAGING:

The device is supplied in Polypropylene case contained within a heat-sealed Tyvek sterilizable peel pouch, along with IFU, inner labels in a duplex box with external identification labels.

Implant Card and Traceability

An implant card is supplied with the device in accordance with MDR Article 18.

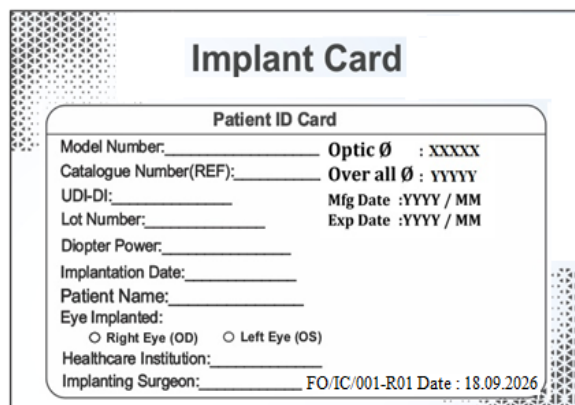
The healthcare institution or surgeon shall complete the implant card with patient identification, device name, model, lot number, UDI, manufacturer details, implantation date, and healthcare institution details.

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Peel-off labels should be affixed to the patient medical record, implant card, and hospital traceability record as applicable.

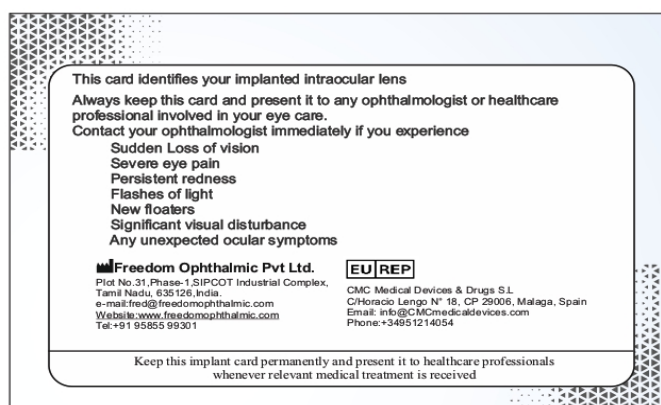
The patient should be instructed to retain the implant card and present it to healthcare professionals during future ophthalmic care or relevant medical



Implant Card

Patient ID Card

Model Number: _____ Optic Ø : XXXXX
 Catalogue Number(REF): _____ Over all Ø : YYYYY
 UDI-DI: _____ Mfg Date :YYYY / MM
 Lot Number: _____ Exp Date :YYYY / MM
 Diopter Power: _____
 Implantation Date: _____
 Patient Name: _____
 Eye Implanted:
☐ Right Eye (OD) ☐ Left Eye (OS)
 Healthcare Institution: _____
 Implanting Surgeon: _____ FO/IC/001-R01 Date : 18.09.2026



This card identifies your implanted intraocular lens
 Always keep this card and present it to any ophthalmologist or healthcare professional involved in your eye care.
 Contact your ophthalmologist immediately if you experience

- Sudden Loss of vision
- Severe eye pain
- Persistent redness
- Flashes of light
- New floaters
- Significant visual disturbance
- Any unexpected ocular symptoms

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 Email: info@CMCmedicaldevices.com
 Phone: +34951214054

Keep this implant card permanently and present it to healthcare professionals whenever relevant medical treatment is received

NOTICE FOR USER / OR PATIENT:

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the User and/or Patient is established.

The instructions provided above have been validated by the manufacturer of the medical device as being Capable of preparing a medical device for safe use.

Equipment & instrument for implantation of IOLs depends on the operating procedures.

STERILIZATION & EXPIRY DATE:

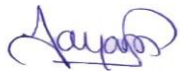
The device is sterilized by Ethylene Oxide (EO) & 60 months from the date of manufacturing

WARRANTY:

We warrant that reasonable care has been used in the design and manufacture of IOL. This warranty is limited to replacement of IOL only. We shall not be liable for any incidental or consequential loss, damage, or expense directly or indirectly arising from the use of this device. We do not assume or authorize any other person to assume on our behalf, any other or additional liability or responsibility in connection with use of this device.

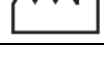
ELECTRONIC VERSION OF IFU:

Available on the website www.freedomophthalmic.com

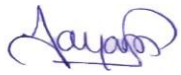
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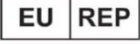
EXPLANATION OF SYMBOLS:

Symbol	Meaning	Symbol	Meaning
	Lot Number/ Batch Number		Sterilization by Ethylene Oxide
	Warning see instruction		Use Until (YYYY-MM)
	Don't use when packing damaged		Humidity Limitation
	Do not reuse		Do not re-sterilize
	Keep away from sunlight		Keep dry
	Temperature limit		Manufacturer
	Country of Manufacturer		CE Mark
	Single Sterile Barrier System		Medical Device
	Patient number		Unique Device Identifier
	Date of Manufacture		Consult instructions for use or Consult electronic instructions for use
	Catalogue number		Country Code
	Authorized representative in European Union		

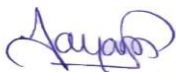

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