

INSTRUCTION FOR USE

Core Surgical Ophthalmic Knife

Content

Each blister contains a single sterile ophthalmic knife. The blades are manufactured from medical grade flat stock stainless steel. The handles are manufactured from medical grade co-polymer (polycarbonate). The handles are coloured to differentiate between distinct specifications of eye knife.

Set Colour	Knife Type
Lilac	Main incision
White	Sideport / Paracentesis

Intended Purpose

The intended purpose of Core Surgical's ophthalmic knife range is to create an incision(s) of a specific size as required during routine ophthalmic surgical procedures.

Indications For Use

Core Surgical's ophthalmic knife range is used by qualified ophthalmic surgeons during routine ophthalmic surgical procedures to enable subsequent treatment of any patient (no specific demographic) with conditions such as cataract and glaucoma.

The choice of ophthalmic knife for a particular procedure is at the discretion of the qualified ophthalmic surgeon.

Contraindications

No known contraindications

Shelf Life

5 years

UKCA Marked

Warnings / Precautions

- The Core Surgical Ophthalmic Knife is designed for single use and must not be re-sterilised to avoid risk of infection.
- Do not use the instrument after expiry date or if the packaging is damaged as device sterility and performance may be compromised.
- Prior to use, inspect the Ophthalmic Knife. Do not use if the blade or handle are damaged.
- The Core Surgical Ophthalmic Knife is only to be used by qualified professionals.
- Used sharps may be contaminated. Disease may occur from injury.
- Used knives should be disposed of in puncture-resistant container.
- The manufacturer shall not be liable for any injury or damage suffered by a patient due to the misuse of the product.

Explanation of Packaging Symbols

	Single use only		Article number
	Do not use after expiry date		Sterilisation by γ-irradiation
	Batch / Lot number		Identification of Legal Manufacturer
	Do not re-sterilise		UKCA Mark including Approved Body ID number
	Single Sterile Barrier System		Sterile barrier system with protective packaging — outer packaging (non-sterile) protects the sterile barrier
	Medical Device		Consult Electronic Instructions For use accompanied by website location
	Caution: take care when handling device		Indicates that product should not be used if the package has been damaged or opened
	Indicates a carrier that contains Unique Device Identifier information		Device manufactured in the United Kingdom (GB)
	Keep Dry		