## CORONET

## **Long Handled Trephine Specification Sheet**





#### **Product Overview**

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The CORONET® Long Handled Trephine has been specifically designed to deliver a reliable and consistent vertical incision, giving improved wound architecture and reduced endothelial cell loss.

The trephine features exclusive CORONET® *Cathedral* Blade Technology, creating an ultra-thin and sharp blade, to minimise undercut and tissue distortion.

- Ultra-sharp and thin blade provides improved approximation between the donor button and patient cornea
- Constructed with hollow core window to increase visualisation of surgical site
- All metal constructed blade with non-glare finish
- Ergonomically balanced handle to increase stability and control
- Double sided Cathedral blade ensures the graft button is not retained once cut
- Compatible with CORONET® Artificial Anterior Chamber (51-935)
- Supplied sterile, single use only, one per box, declared 5-year shelf life
- Manufactured in the UK



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## **Long Handled Trephine Size Options**

Size (mm)	51-903
6.00	51-903-6.00
6.25	51-903-6.25
6.50	51-903-6.50
6.75	51-903-6.75
7.00	51-903-7.00
7.25	51-903-7.25
7.50	51-903-7.50
7.75	51-903-7.75
8.00	51-903-8.00
8.25	51-903-8.25
8.50	51-903-8.50
8.75	51-903-8.75
9.00	51-903-9.00
9.25	51-903-9.25
9.50	51-903-9.50
9.75	51-903-9.75
10.00	51-903-10.00

### **Material Specification**





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Product Component	Specification
Trephine blade	Stainless Steel
Trephine Body	Aluminium
Pouch	Tyvek/Film
Outer Box/Carton	500 Micron Printed White Boxboard

#### **Intended Use**

The CORONET® Long Handled trephine is to be used for either trephination of the donor button or trephination of the recipient eye during corneal transplant surgery. The ultra-thin blade profile cuts cleanly through the corneal tissue allowing for a better approximation between the donor button and the patient cornea.

#### **Instructions for Use**

The instructions for use and suggested surgical technique are supplied in the form of a multi-lingual leaflet with instructions given in diagram form where appropriate. The international symbols used on the packaging are explained in each language. A leaflet is supplied with each product

### **Sterilisation**

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Products are sterilised by Ethylene Oxide (EO) according to a validated cycle and the requirements of BS EN ISO 11135-1 (current).

### **Conformity to the European Directives**

The CORONET® range of products are classified as a surgically invasive device intended for transient use, (Rule 6, Annex IX 93/42/EEC Medical Devices Directive, as amended by 2007/47/EEC) and is therefore classified as a Class IIa device.

