

BIOMEDICAL ENGINEERING DEPARTMENT
ADDU EQUATORIAL HOSPITAL
TECHNICAL SPECIFICATION
DISPOSABLE STERILE LASER FIBRES FOR LIMMER DIOLAS LASER SYSTEM

General Requirements

1. The offered laser fibres shall be new, original, sterile, single-use disposable products.
2. The fibres shall be fully compatible with the hospital's existing **Limmer DIOLAS diode laser system** without requiring any additional adapters, connectors or hardware modifications
3. The offered fibres shall be supplied with the manufacturer's original packaging and labeling.
4. All fibres shall be individually packed and sterilized.
5. Sterility status and expiry date shall be clearly indicated on the packaging.
6. The bidder shall submit the manufacturer's catalogue reference (part number) for the offered item.
7. Product shall have CE Mark, FDA clearance, or equivalent internationally recognized regulatory approval.
8. The supplier shall provide a written declaration confirming compatibility with the existing Limmer DIOLAS laser system.

Item 1

Sterile Endoradial Fibre – 400 μm

Reference Product

Limmer Part Number: L10.04.100003

Technical Requirements

1. Sterile, single-use disposable radial laser fibre.
2. Fibre core diameter: **400 μm** .
3. Radial laser energy emission.
4. Intended for endovenous laser ablation and minimally invasive laser procedures.
5. Flexible fibre construction suitable for clinical use.
6. Compatible with the existing **Limmer DIOLAS diode laser system**.
7. Supplied sterile and ready for immediate use.
8. Manufacturer catalogue reference shall be **L10.04.100003** or equivalent fully compatible product.

