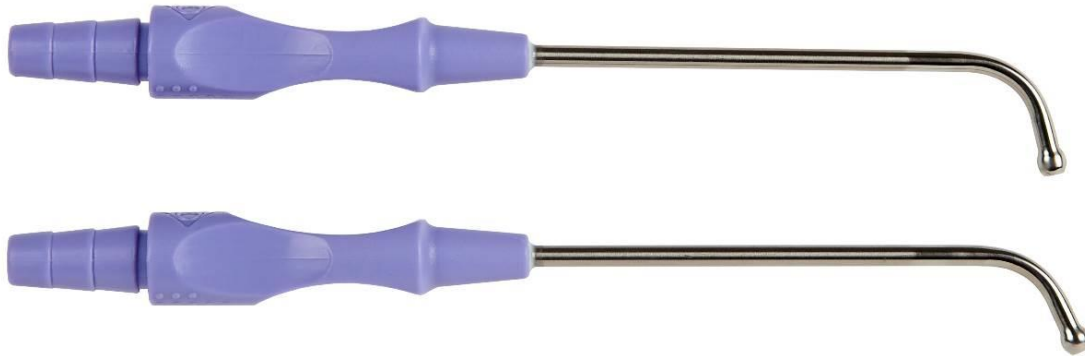


F.E.S.S Suction Tube Range Specification Sheet



Product Overview

The NETWORK ENT® F.E.S.S Suction Tube range has been specifically designed to provide precision and control during Functional Endoscopic Sinus Surgery (F.E.S.S). The olive shaped suction tip permits atraumatic entrance into the sinus ostium, so that suction dissection or removal of synechiae can be performed.

Each suction tube is manufactured without bending or pinching to create a continuous radius, significantly reducing the risk of blockage.

- Constructed from high quality Japanese steel for superior thin walled tubing
- Curved tip for maximum visibility and access to difficult areas
- Ergonomically balanced and lightweight handle for unrivalled manoeuvrability and control
- Supplied sterile, single use only, declared 5 year shelf life
- Manufactured in the UK

F.E.S.S Suction Tube Size Options

Product Size	Box Quantity	Product Code
Sinus suction tube, 3mm x 85mm, distal bend, 80 degree angle with olive tip c/w control hole	25	74-2060
Sinus suction tube, 3mm x 85mm, distal bend, 65 degree angle with olive tip c/w control hole	25	74-2063

F.E.S.S Suction Tube Range Specification Sheet

Material Specification

Product Component	Specification
Tube	Stainless Steel
Moulded handle	Polypropylene (PP)
Pouch Packaging	Tyvek/Film
Outer Box/Carton Packaging	500 Micron White Boxboard

Intended Use

The NETWORK ENT® F.E.S.S Suction tubes are designed specifically for ENT procedures and are intended for the removal of excess fluids and debris from the surgical site.

Sterilisation

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

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.

Conformity to the European Directives

The Network range of Aspiration Suction Tubes and Fine Ends are identified as surgically invasive devices for transient use and are therefore classified as a Class IIa devices according to Annex IX, Rule 6, of the Medical Devices Directive 93/42/EEC as amended by 2007/47/EC.