NETWORK ENT PRODUCTS

Suction Sets Range Specification Sheet

Product Overview

The NETWORK ENT® Suction Sets range, has been specifically designed to provide precision and control during ENT procedures.

- Complete suction system includes suction tube with handle, tubing and adapter
- Compatible with all commercially available suction systems
- Supplied sterile, single use only, declared 5 year shelf life
- Manufactured in the UK

Suction Sets Range Size Options

Product Size	Box Quantity	Product Code
Suction set complete with house adapter, with suction control hole	50	74-2003
Suction set with Zoellner type 2mm x 85mm, 10 degree shallow distil bend (2.0mm/6Fr)	50	74-2004
Suction set with Zoellner type 2mm x 85mm, 30 degree acute distal bend (2.0mm/6Fr)	50	74-2005
Suction set with Belucci type 2mm x 85mm, proximal bend, with suction control (2.0mm/6Fr)	50	74-2006

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Material Specification

Product Component	Specification
Tube (74-2004,74-2005 & 74-2006 only)	Stainless Steel
Moulded handle	Polypropylene (PP)
House Adapter (74-2003 only)	ABS
Reducer connector	Polyamide
Connector Tubing	Silicone
Pouch Packaging	Tyvek/Film
Outer Box/Carton Packaging	500 Micron Brown Boxboard

Intended Use

The NETWORK ENT® Suction Sets 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.

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