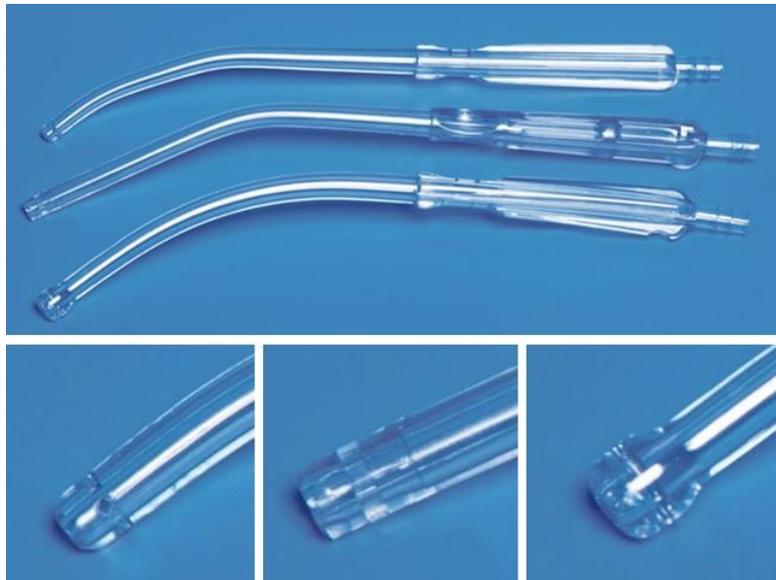


## Yankauer Suction Tube Range Specification Sheet



### Product Overview

The NETWORK ENT® Yankauer Suction Tube range has been specifically designed to quickly and safely clear operative sites during a procedure.

- Manufactured with clear tips for improve visibility
- Ribbed universal connector allows secure fit to all types of tubing
- Ergonomically balanced handle for improved precision
- Available in a variety of type, size and angles
- Supplied sterile, single use only, declared 5 year shelf life

## Yankauer Suction Tube Range Specification Sheet

### Yankauer Suction Tube Range Size Options

Product Size	Box Quantity	Product Code
8mm rigid open tip, small curve, with suction control	50	74-1211
8mm rigid open tip, small curve, without suction control	50	74-1212
6mm rigid paediatric tip, medium curve, with suction control	50	74-1213
6mm rigid paediatric tip, medium curve, without suction control	50	74-1214
10mm rigid bulbous tip, large curve, with suction control	50	74-1215

### Material Specification

Product Component	Specification
Yankauer handle	k-resin
Pouch Packaging	Arjor and film pouch

### 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

## Yankauer Suction Tube Range Specification Sheet

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.