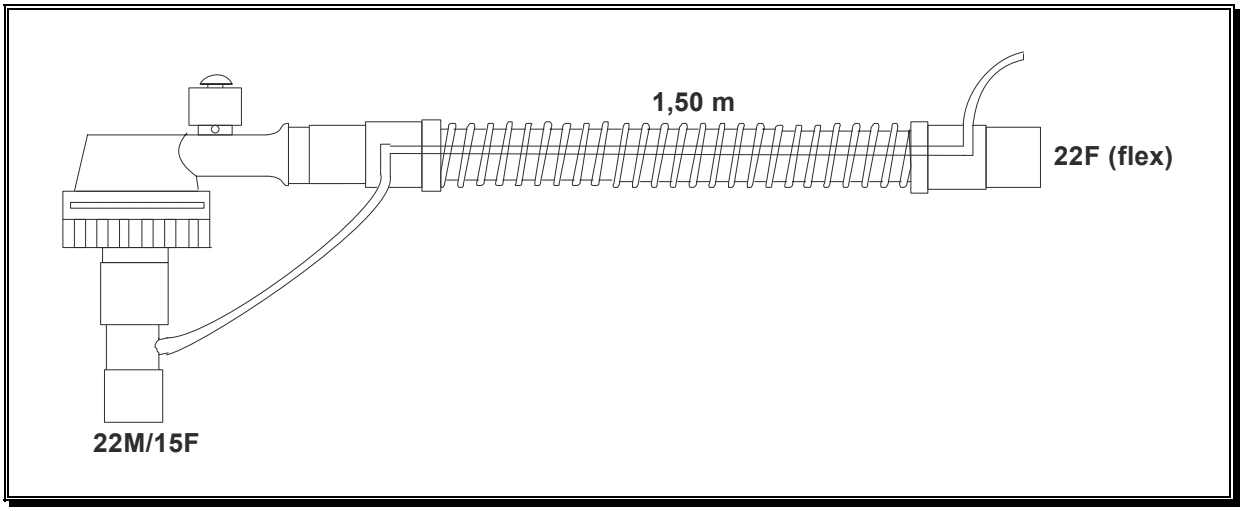
TECHNICAL DATA SHEET

REF: 02940



Description and application:	Breathing circuit for SIEM ventilators type BA 2001 XX – EL.
Length:	1.50 m.
Features:	• Transparent, extremely flexible and light tubing, Ø 19 mm, smoothbore inside to avoid water stagnation and consequent bacterial growth, with rigid spiral to avoid any likelihood of lumen reductions caused by kinking. • Translucent patient valve with a built-in 22M/15F 360° patient swivel port to connect masks or tracheal tubes/tracheotomy cannulas. • Equipped with a 60 cmH_2O pressure relief valve to reduce possibility of lung damage. • Pressure monitoring tube inserted inside the ventilation tube for greater ease of use and minimal bulk. • 22F flexible and soft machine-side connector for a perfect seal.
Compliance:	On a length of 1 m. = $ml \cdot kPa^{-1} \cdot m^{-1}$ 4,6 or = $ml \cdot cmH_2O^{-1} \cdot m^{-1}$ 0,46. (tests carried out at a pressure of 6 kPa and in accordance with EN 12342)

DOCUMENT REFERENCES:		
DRAWN UP BY: Quality Group	APPROVED BY: (DOMENICO SCARDOVI)	DATE: 25/09/2025
File name: 02940-1e.doc	Rev.: 01	Page/Tot. Pages.: 1/2

Sterilisation type: ***(product available clean or sterile packaged)***	Sterilized with Ethylene Oxyde (ETO) under a process validated in accordance with EN EN 11135-1 standard. Desorbing process to ensure ETO residual limits assessed by ISO 10993-7. Certificates available on request.
Raw materials:	PVC without phthalates (DOP/DEHP free); PC. Phthalates (DEHP/DOP) and Latex free
Packaging:	STERILE PRODUCT (individually packaged): PRIMARY: pouch (Medical Kraft Paper, Film PA/PE) conforming to EN ISO 11607-1 and EN ISO 11607-2; SECONDARY: Carton boxes. NON-STERILE PRODUCT (individually packaged): PRIMARY: PE bags; SECONDARY: Carton boxes.
Pieces per box:	REF / STERILE PRODUCT / NON STERILE PRODUCT (add "NS" at the end of the code) 02940 / box of 25 pcs single packed / box of 25 pcs single packed
Storage and disposal:	The device must be stored respecting the instructions given by the symbols on the packaging. After use, the device must be treated and/or disposed of as hazardous medical waste according to local regulations in force.
CE certification details :	Class **IIa** medical device; the device is certified by the Notified Body TÜV SÜD Product Service GmbH - Ridlerstrasse 65 - 80339 Munich, CE0123 according to the procedure of Annex V of Directive 93/42/EEC as amended, in accordance with the transitional provisions of Article 120.2 of Regulation (EU) 2017/745.
Manufacturer: *in accordance with Directive 93/42 EC of 14/06/93*	