

# Terumo® Surflo® Winged Infusion Set

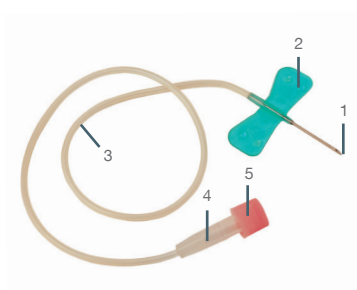
## Product Information Card



### Description:

The Surflo® winged infusion set (= Scalp Vein or SV-set) is intended to introduce or withdraw fluids from the vascular system after vein puncture.

### Components:



Description	Raw Materials
1. Cannula	Stainless steel
2. Wing hub	Polyvinyl chloride + master batch
3. Tube	Polyvinyl chloride
4. Adapter: BL item code EL/BLK item code	Polypropylene PMMA
5. Adapter cap: BL item codes EL/BLK item code	Polyethylene + master batch HDPE + master batch
Protector	Polyethylene



### Product Range:

Product Code	Colour	External Cannula Ø	Cannula Length	Dead Space Volume	Tubing Length
SV*18BLK03	Pink	18G 1.20 mm	19 mm	0.46 ml	300 mm
SV-19BL	Ivory	19G 1.10 mm	19 mm	0.43 ml	300 mm
SV-21BL	Green	21G 0.80 mm	19 mm	0.42 ml	300 mm
SV-22BL	Black	22G 0.70 mm	19 mm	0.42 ml	300 mm
SV-23BL	Blue	23G 0.60 mm	19 mm	0.41 ml	300 mm
SV-25BL	Orange	25G 0.50 mm	19 mm	0.41 ml	300 mm
SV*27EL	Grey	27G 0.40 mm	19 mm	0.27 ml	200 mm

# Terumo® Surflo® Winged Infusion Set

## Packaging:

**Single unit:** blister pack (thermoformed blister film)

Product Code	Unit Box	Shipping Carton	Gross Weight	Dimensions (LxWxH)
SV*18BLK03	50	500	3.2 kg	430 x 220 x 265 mm
SV-19BL	50	500	2.6 kg	438 x 228 x 276 mm
SV-21BL	50	500	2.6 kg	438 x 228 x 276 mm
SV-22BL	50	500	2.6 kg	438 x 228 x 276 mm
SV-23BL	50	500	2.6 kg	438 x 228 x 276 mm
SV-25BL	50	500	2.6 kg	438 x 228 x 276 mm
SV*27EL	100	1000	n.a.	475 x 255 x 465 mm

## Sterilization:

With ethylene oxide to a Sterility Assurance Level of at least 10<sup>-6</sup> for all BL/EL-items  
With gamma radiation to a Sterility Assurance Level of at least 10<sup>-6</sup> for BLK-item

## Storage:

Shelf life of 5 years for all BL-items / 3 years for all EL/BLK-items

\* The lot number and expiry date are embossed in the film of the thermoformed blister pack

\* Do not store at extreme temperature and humidity

## Precautions for Use:

Check integrity of the unit package before use  
For single use only  
Dispose of safely after single use to avoid risk of infection

## Status of Regulation:

Class IIa, medical device following MDD 93/42/EEC

Notified body: TÜV Rheinland 0197  
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