Suction Coagulation Probe

Used for suction and coagulate during endoscopic mucosal resection

Art. No.	Tip Ø (mm)	Sheath Ø (mm)	Working Length (cm)	Working Channel Ø (mm)
SCP	2.5	2.6	230	2.8

Class IIb device – for single use only

Wound Drainage and Suction

Silicone Bulb 100ml
Drainage feature with a radiopaque stripe for x-ray detection.

Art. No.	Productname	Size (Fr/CH)	Length (mm)
CDD12	Silicone bulb and tube	12	900
CDD14	Silicone bulb and tube	14	900
CDD16	Silicone bulb and tube	16	900
CDD18	Silicone bulb and tube	18	900
SPS 12	Trocar only		
SPS 14	Trocar only		
SPS 16	Trocar only		
SPS 18	Trocar only		

Class IIa device – for single use only

Multiband Ligator

Barrel with 6 bands

Used for endoscopic ligation of esophageal varices at or above gastroesophageal junction

Fits on wide range of endoscopes
Second last band light colour
Barrel with 7 bands available

Standard Band Ligator
Requires 9.5 to 11.2 mm
Endoscope diameter

Paediatric Band Ligator
Requires 8.0 to 9.5 mm
Endoscope diameter

	Art. No.	Description
Standard	**SGK-6**	Set incl. Applicator barrel, thread puller and flushing needle
	SGB-6	Barrel and thread puller only
Paediatric	**SGK-6-S**	Set incl. Applicator barrel, thread puller and flushing needle
	SGB-6-S	Barrel and thread puller only

Class IIa device – non sterile – for single use only

Multiband Ligator Colonic

Used for endoscopic ligation of colonic varices

Fits on wide range of endoscopes
Second last band light colour

Band Ligator Colonic Set contains:
Applicator, Barrel loaded with 6 bands
or Barrel loaded with 7 bands
Atraumatic plastic hook with tube
Flushing needle blunt tip
Second last band with light colour
Requires 12.8 to 13.5 mm
Endoscope diameter

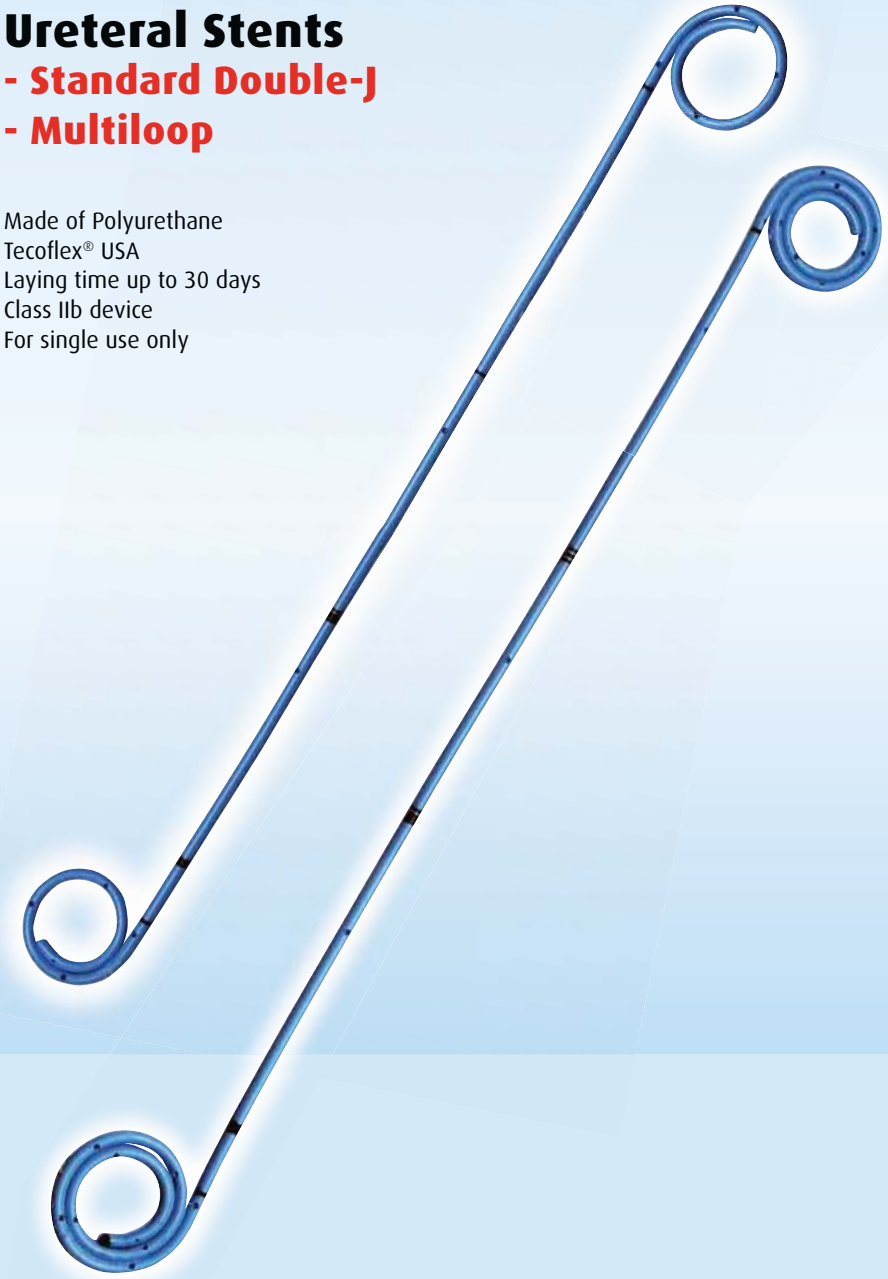
Barrel Colonic contains:
Barrel loaded with 6 bands
or Barrel loaded with 7 bands
Atraumatic plastic hook with tube

	Art. No.	Description
Band Ligator	**SBLS-6-C**	Multiband Ligator Colonic Set incl. Applicator barrel, thread puller and flushing needle
	SBL-6-C	Multiband Ligator Colonic Barrel and thread puller only
Barrel Colonic	**SBLS-7-C**	Multiband Ligator Colonic Set incl. Applicator barrel, thread puller and flushing needle
	SBL-7-C	Multiband Ligator Colonic Barrel and thread puller only

Class IIa device – non sterile – for single use only

Ureteral Stents
- Standard Double-J
- Multiloop

Made of Polyurethane
Tecoflex® USA
Laying time up to 30 days
Class IIb device
For single use only



MarFlow®

Ureteral Stents
Standard Double-J and Multiloop

Set consists of stent, pusher, suture and clamp.

One side open	Both sides open			
Art. No.	Art.No.	Size (Fr/CH)	Length (cm)	Guidewire
SS-4716	**SOT-4716**	4.7	16	0.032″
SS-4724	**SOT-4724**	4.7	24	0.032″
SS-4726	**SOT-4726**	4.7	26	0.032″
SS-4728	**SOT-4728**	4.7	28	0.032″
SS-4730	**SOT-4730**	4.7	30	0.032″
SS-47ML+ML	**SOT-47ML+ML**	4.7	Multiloop	0.032″
SS-516	**SOT-516**	5	16	0.035″
SS-524	**SOT-524**	5	24	0.035″
SS-526	**SOT-526**	5	26	0.035″
SS-528	**SOT-528**	5	28	0.035″
SS-530	**SOT-530**	5	30	0.035″
SS-5ML+ML	**SOT-5ML+ML**	5	Multiloop	0.035″
SS-616	**SOT-616**	6	16	0.035″
SS-624	**SOT-624**	6	24	0.035″
SS-626	**SOT-626**	6	26	0.035″
SS-628	**SOT-628**	6	28	0.035″
SS-630	**SOT-630**	6	30	0.035″
SS-6ML+ML	**SOT-6ML+ML**	6	Multiloop	0.035″
SS-716	**SOT-716**	7	16	0.038″
SS-724	**SOT-724**	7	24	0.038″
SS-726	**SOT-726**	7	26	0.038″
SS-728	**SOT-728**	7	28	0.038″
SS-730	**SOT-730**	7	30	0.038″
SS-7ML+ML	**SOT-7ML+ML**	7	Multiloop	0.038″
SS-826	**SOT-826**	8	26	0.038″
SS-828	**SOT-828**	8	28	0.038″
SS-830	**SOT-830**	8	30	0.038″
SS-8ML+ML	**SOT-8ML+ML**	8	Multiloop	0.038″

other sizes available

Set with stainless steel guidewire add +G (for example SS-4716+G).
Set with nitinol guidewire add +GTHS (for example SS-4716+GTHS).

3EC International a.s., Hraničná 18, 821 05 Bratislava, Slovakia
Notified Body No. 2265

EC CERTIFICATE

No. 2013-MDD/QS-032

issued in compliance with the Council Directive 93/42/EEC as amended by 2007/47/EC, which is implemented by the Slovak Government Decree No. 582/2008 Coll. certifies that the medical device of Class IIa & IIb,

Medical Devices for Gastroenterology
(for detailed list refer to Annex, pages 1 to 3)

manufactured by company

Marflow AG
Soodstrasse 57, CH-8134 Adliswil, Zürich, Switzerland

is manufactured under conditions fulfilling the quality system requirements of Annex II, excluding (4), of the Directive 93/42/EEC as amended by 2007/47/EC.

The Notified Body No. 2265 has performed an audit of the above device quality system. The full quality assurance system has been assessed and found that it meets the requirements above. The quality system is subject to continuous surveillance according to Annex II, Sections 3.3, and 5, of the Directive 93/42/EEC as amended 2007/47/EC. The detailed description of the system, requirements and measures applied by the manufacturer are presented in the Audit Reports No. 310121A and 310121B, and the Final protocol No. 310121b/2013 that is enclosed to this certificate.

This certificate is issued under the following conditions:

It applies only to the quality system maintained in the manufacture of the above referenced model of medical device and it does not substitute the design or type-examination procedures, if requested. The certificate remains valid until the manufacturing conditions or the quality system are changed but until August 26th, 2019 at the latest. The certificate validity is conditional upon positive results of regular surveillance audits and fulfilment of relevant legal and other requirements by manufacturer.

In Bratislava, on August 27th, 2013

Notified body 3EC International NB 2265 NOTIFIKOVANÁ OSOBA

Dr. Katarína Srdošová
Responsible to act on behalf of NB 2265

TRANSGLOBAL QUALITY ASSESSORS LLP

Management System Certificate

Certificate No. MD.QMS.91.006.06.16

This is certify that

Marflow AG

at

Soodstrasse 57, CH-8134 Adliswil, Zürich, Switzerland.

has been found to conform to Management System Standard

ISO 13485: 2003

This certificate valid for the following product / service ranges:

DESIGN AND MANUFACTURE OF NON-ACTIVE AND ACTIVE (NON-IMPLANTABLE) MEDICAL DEVICES FOR UROLOGY AND GASTROENTEROLOGY

Internal Certification : 23.06.2016
Valid until : 22.06.2019

(Authorized Signatory)
Transglobal Quality Assessors LLP

This is an accredited certificate authorised for issue by Accreditation Services for certifying bodies (Europe) Limited, who have assessed Transglobal Quality Assessors LLP Located at PUNE, INDIA, against defined criteria and in cognisance of ISO 17021, "Conformity Assessment Requirements for bodies providing audit and certification of management systems". This certificate is only valid when confirmed by register listed in the International register of Quality Assessed Organisation : www.irqao.com