ISOLATED LESION MEASUREMENT/ DEVICE SELECTION FORM



Confidential patient information — Do not disclose legally protected data.

The following information	on is required to ensure that	the appropriate devices an	id backups are available	e for the procedure

atier	nt ID:						Institution:	
hysi	cian:						Imaging date:	
уре о	of Aneui	rysm,	/lesion:					
			Location		Measurement List single value used to select devices	Range List range of measurement taken	CT table position/angio Specify CT frame number or specify angio	NOTES
			DIAMETER					
ΙΛ	$\wedge \wedge 1$	— А	Proximal implantation si	te	mm	mm		
		В	1 cm from proximal implantation site		mm	mm		
سمر		C	2 cm from proximal implantation site		mm	mm		
		D	Maximum aneurysm/lesi	ion	mm	mm		
		E	2 cm from distal implantation site		mm	mm		
VV	\mathcal{N}	F	1 cm from distal implantation site		mm	mm		
VV	<u> </u>	−G	Distal implantation site		mm	mm		
		Н	Right common iliac		mm	mm		
		1	Left common iliac		mm	mm		
		J	Right extension iliac/ femoral		mm	mm		
		К	Left extension iliac/femo	oral	mm	mm		
LENG								
L¹	Proxi Dista		neck rom aneurysm/lesion					
			clavian		cm	cm		1
L ² Proximal neck Distance from aneurysm/lesion to left common carotid artery			cm	cm				
M Aneurysm/lesion Length of aneurysm/lesion segment			cm	cm		7/1		
N	Dista	l nec						
	Dista to cel		om aneurysm/lesion		cm	cm		SUGGESTED C-ARM ANGLE
0			ment length		cm	cm		RAO
ANG								LAO
Р	Proxi	mal a	ingle		o			LAU
Q	Dista	l ang	le (if applicable)		0			LATERAI
	nere sign lantatior		t calcium/thrombus at the?	e pro	oximal	YES	□ №	NOTES
	nere sign lantatior		t calcium/thrombus at the	e dis	tal	☐ YES	□ №	
Is treatment length 10 cm or less?				☐ YES	□ NO			
If yes, will both necks (proximal and distal) accommodate a single device?				YES	□ №			
Is there a plan for coverage of the left subclavian?			n?	YES	□ NO			
If yes, is transposition or bypass clinically indicated				idicated?	YES	□ №		
Is angle less than 60°?				☐ YES	□ №			
If yes, is neck length greater than 2 cm?				YES	□ №	(See reverse for device selection form		

The following information is require		ate devices	and any additional devi	ces are available for th	e procedure.
Patient ID:			Institution:		
Physician:			Imaging date:		
Intended device introduction site	Right Left	☐ Iliac ☐ Femo		enal aorta 🔲	Conduit
TREATMENT OPTION 1					
Devices listed as implanted (proximal to distal)	Order of implantation (#1, #2, etc.)		Intended aortic diameters (mm)	Labeled diameter (mm)	Device length (cm)
			16-19.5	21	10
		[19.5-24	26	10
		[22-26	28	10, 15
		[24–29	31	10, 15, 20
TREATMENT OPTION 2			27–32	34	10, 15, 20
Devices listed as implanted (proximal to distal)	Order of implantation (#1, #2, etc.)		29-34	37	10, 15, 20
(proximar to distar)	(#1, #2, 666.)		31–37	40	10, 15, 20
		-	34-42	45	10, 15, 20

Intended GORE® TAG® Conformable Thoracic Stent Graft size: (Check all device sizes and indicate number of each size to be ordered.)

Device size (mm x cm)	QTY.	Catalogue number*	Device size (mm x cm)	QTY.	Catalogue number*	Device size (mm x cm)	QTY.	Catalogue number*
□ 21 x 10		TGM212110	34 (proximal), 28 (distal) x 15		TGM342815			
☐ 26 (proximal), 21 (distal) x 10		TGM262110	☐ 37 (proximal), 31 (distal) x 15		TGMR373115			
□ 26 x 10		TGM262610	☐ 40 (proximal), 34 (distal) x 15		TGMR403415			
☐ 31 (proximal), 26 (distal) x 10		TGMR312610	☐ 45 (proximal), 37 (distal) x 15		TGM453715			
□ 28 x 10		TGM282810	□ 28 x 15		TGM282815			
□ 31 x 10		TGMR313110	□ 31 x 15		TGMR313115	□ 31 x 20		TGMR313120
□ 34 x 10		TGM343410	□ 34 x 15		TGM343415	□ 34 x 20		TGM343420
□ 37 x 10		TGMR373710	☐ 37 x 15		TGMR373715	☐ 37 x 20		TGMR373720
☐ 40 x 10		TGMR404010	☐ 40 x 15		TGMR404015	☐ 40 x 20		TGMR404020
☐ 45 x 10		TGM454510	☐ 45 x 15		TGM454515	☐ 45 x 20		TGM454520

^{*}Some sizes not available in all regions.

GORE® DRYSEAL Flex Introducer Sheath: (outer diameter)

The second secon						
Sheath size (Fr)	Device diameter (mm)	Sheath working length (cm)	QTY.	Catalogue number*		
18 (6.7 mm)	21	33		GDSF1833		
18 (6.7 mm)	21	65		GDSF1865		
20 (7.5 mm)	26-31	33		GDSF2033		
20 (7.5 mm)	26-31	65		GDSF2065		
22 (8.2 mm)	34-40	33		GDSF2233		
22 (8.2 mm)	34-40	65		GDSF2265		
24 (8.8 mm)	45	33		GDSF2433		
24 (8.8 mm)	45	65		GDSF2465		
26 (9.5 mm)		33		GDSF2633		
26 (9.5 mm)		65		GDSF2665		

^{*}GDSF catalogue numbers are not available in all regions; use DSF catalogue numbers instead where appropriate.

GORE® Tri-Lobe Balloon Catheter:

Device size	QTY.	Catalogue number
☐ Aortic diameters 16–32 mm		BCM1634
Aortic diameters 26–42 mm		BCL2645

If the physician's proposed use of the device is outside the scope of the *Instructions for Use*, Gore is providing this information in response to the physician inquiry. Physician, as a medical professional, is responsible for evaluating the information provided and deciding on the use of the device.

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Consult Instructions Refer to Instructions for Use at eith governedical comfor a com eifu.goremedical.com for a complete description of all applicable indications,

warnings, precautions and contraindications for the markets where this product is available. $R_{\!\! X \, \text{Only}}$

