

Retrievable & Permanent

PROVEN & RELIABLE



OPTEASE[®]

Retrievable Vena Cava Filter

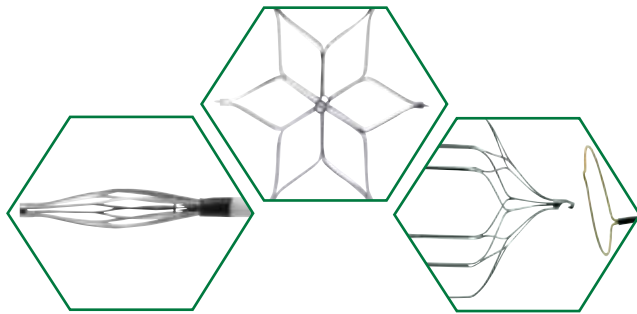
Cordis[®]
A Cardinal Health company



Engineered to Perform. Built to Stay.

OPTEASE® Retrievable Vena Cava Filter

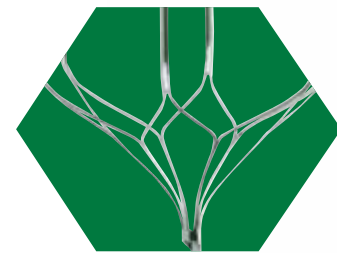
The OPTEASE® Vena Cava Filter was designed to perform predictably and dependably. It allows you to focus on the effective treatment of your patients rather than the risk of caval perforation, migration or strut embolization.



Filter Features

Material	Nitinol
Max Caval Diameter	30 mm
Low Profile	6F Sheath
Ease of Use	Femoral, Jugular and Antecubital Vein Placement Options
Retrieval Direction	Femoral
Retrieval Period (per IFU)	Can be retrieved up to and including 12 days after placement

WARNING: Implant of the OPTEASE® Vena Cava Filter with the hook oriented in the cranial direction can result in life threatening or serious injury including, but not limited to dissection, vessel perforation, migration of the filter with secondary damage to cardiac structures and ineffective pulmonary embolism prevention.



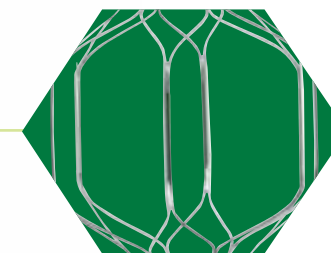
Closed Cage
Designed to eliminate risk of caval perforation and strut embolization



Dual Prong Caudal Hook
For easier snare capture



Fixation Barbs
Reduce migration to maintain clot capture efficiency



Side Struts
Self centering upon insertion minimizing the risk of tilting



Two of the Most Tested Filters on the Market

OPTEASE® Retrievable Vena Cava Filter

PROOF Trial: Protection from Pulmonary Embolism with the Cordis OPTEASE® Retrievable Vena Cava Filter¹

Prospective, multi-center, single filter trial with 150 patients enrolled

PE Protection

Symptomatic PE	
1 Month (n=111)	0.0%
6 Month (n=70)	0.0%

Low Complications

Caval Perforation	
1 Month (n=111)	0.0%
6 Month (n=70)	0.0%
Filter Migration	
1 Month (n=111)	0.9% (1 Case)
6 Month (n=70)	No New Cases
Symptomatic Filter Thrombosis	
1 Month (n=111)	0.8% (1 Case)
6 Month (n=70)	No New Cases

TRAPEASE® Permanent Vena Cava Filter

Cordis TRAPESE® Permanent Vena Cava Filter: Experience in 751 Patients²

The largest single filter, single center VCF with CT follow up

PE Protection

Overall CTPA Proven PE	1.0%
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Low Complications

Migration	0.0%
Caval Penetration	0.0%
Tilting	0.0%
Embolization of Filter Fragments	0.0%
Asymptomatic Near Caval Occlusions	0.7%



OPTEASE®

Retrievable Vena Cava Filter

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Cordis OPTease® Retrievable Vena Cava Filter Ordering Information

Description	Access Site	Catalog Numbers
Cordis OPTease® Vena Cava Filter and Introduction Kit (55 cm)	Femoral	466-F210AF
Cordis OPTease® Vena Cava Filter and Introduction Kit (55 cm)	Jugular	466-F210AJ
Cordis OPTease® Vena Cava Filter and Introduction Kit (90 cm)	Antecubital, Jugular	466-F210BJ

Cordis TRAPEASE® Permanent Vena Cava Filter Ordering Information

Description	Access Site	Catalog Numbers
Cordis TRAPEASE® Vena Cava Filter and Introduction Kit (55 cm)	Jugular, Femoral	466-P306A
Cordis TRAPEASE® Vena Cava Filter and Introduction Kit (90 cm)	Antecubital, Jugular, Femoral	466-P306B

¹Ziegler, JW et al. PROOF Trial: Protection from Pulmonary Embolism with the OPTease® Filter. JVIR 2008; 19:1165-1170.

²Kalva, SP et al. TRAPEASE® Vena Cava Filter: Experience in 751 Patients. J Endovasc Ther 2006; 13:365-372.

WARNING: Implant of the OPTease® Vena Cava Filter with the hook oriented in the cranial direction can result in life threatening or serious injury including, but not limited to dissection, vessel perforation, migration of the filter with secondary damage to cardiac structures and ineffective pulmonary embolism prevention.

IMPORTANT INFORMATION:

Prior to use, refer to the full "Instructions For Use" supplied with these devices for more information on indications, contraindications, suggested procedures, warnings and precautions. Contact your Cordis sales representative for availability and ordering. As part of the Cordis policy of continuous product development, we reserve the right to change product specifications without prior notification. EMEA For Healthcare Professionals only. EU790 2/16.

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