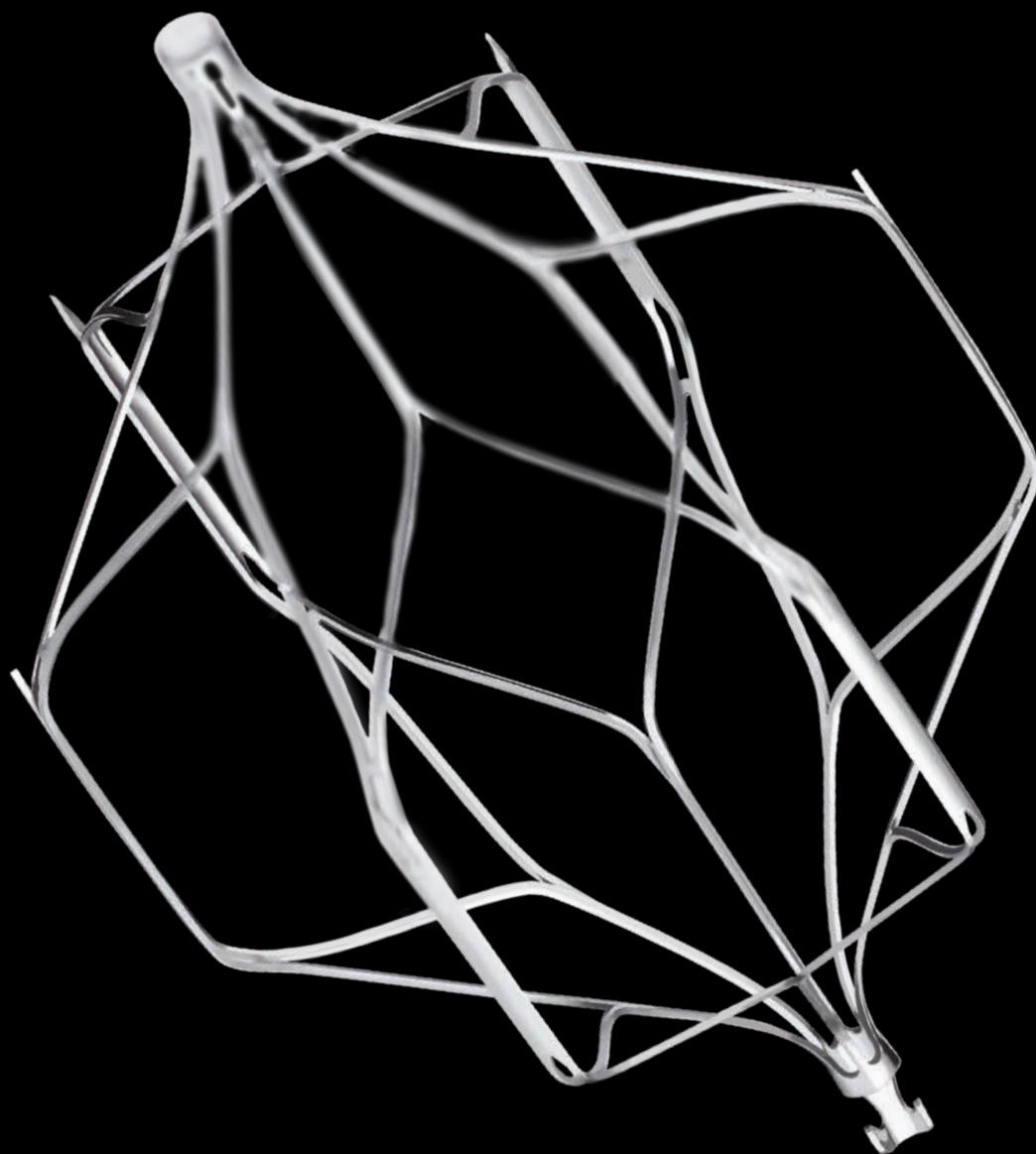


RETRIEVABLE,
PROVEN &
RELIABLE

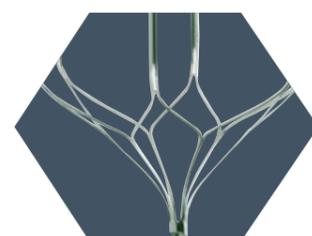
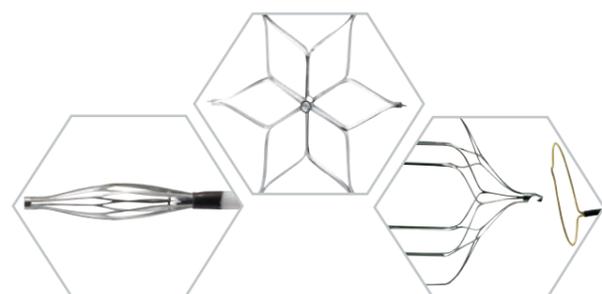


OPTEASE™
RETRIEVABLE VENA CAVA FILTER



Engineered to Perform. Built to Stay.

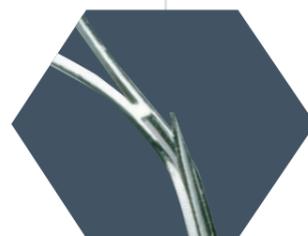
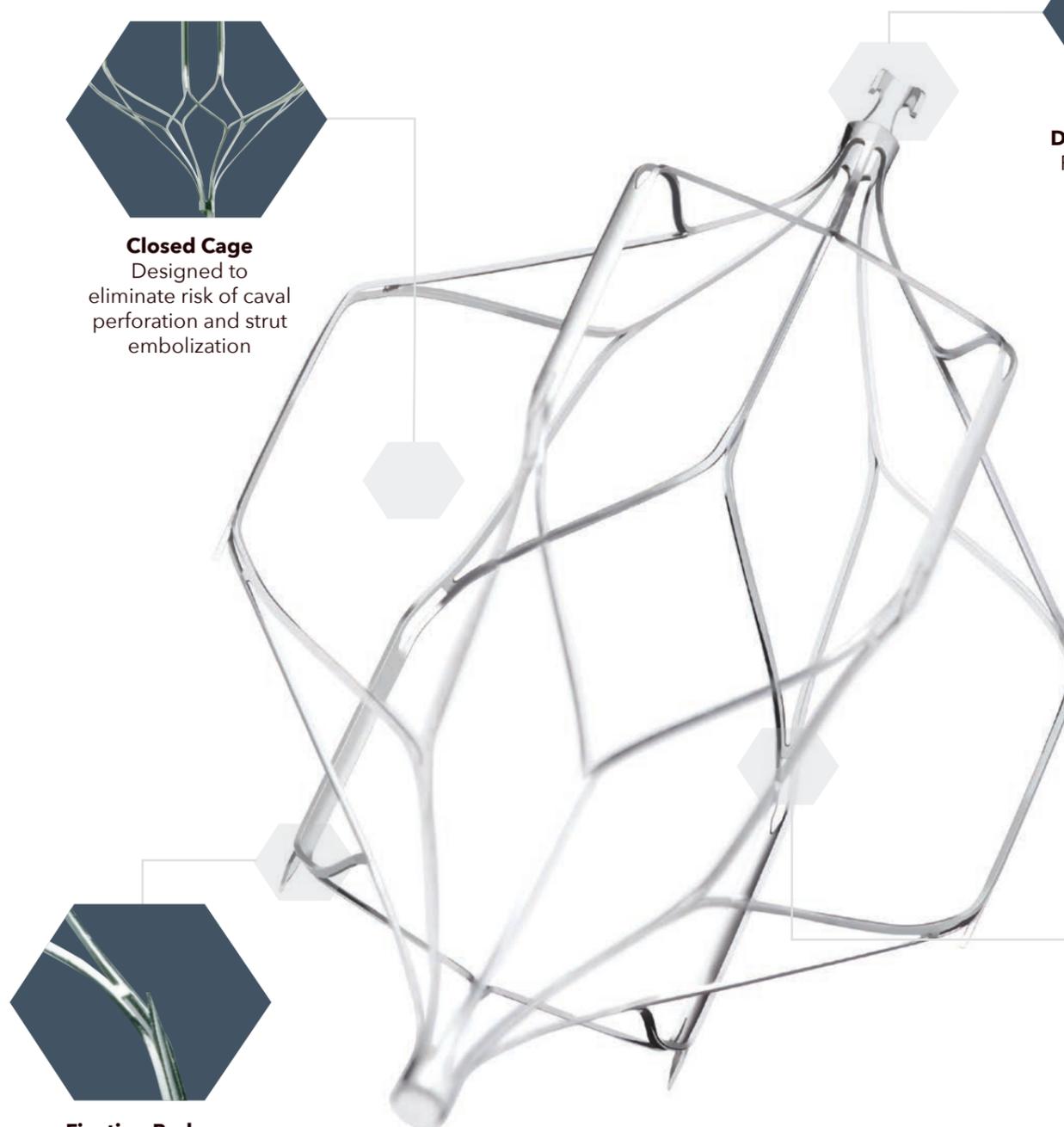
The OPTEASE™ Vena Cava Filter was designed to perform predictably and dependably. It allows you to focus on the effective treatment of your patients rather than the risk of caval perforation, migration or strut embolization.



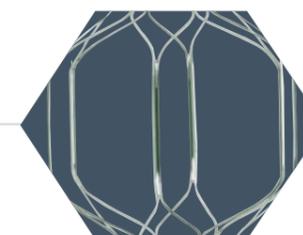
Closed Cage
Designed to eliminate risk of caval perforation and strut embolization



Dual Prong Caudal Hook
For easier snare capture



Fixation Bars
Reduce migration to maintain clot capture efficiency

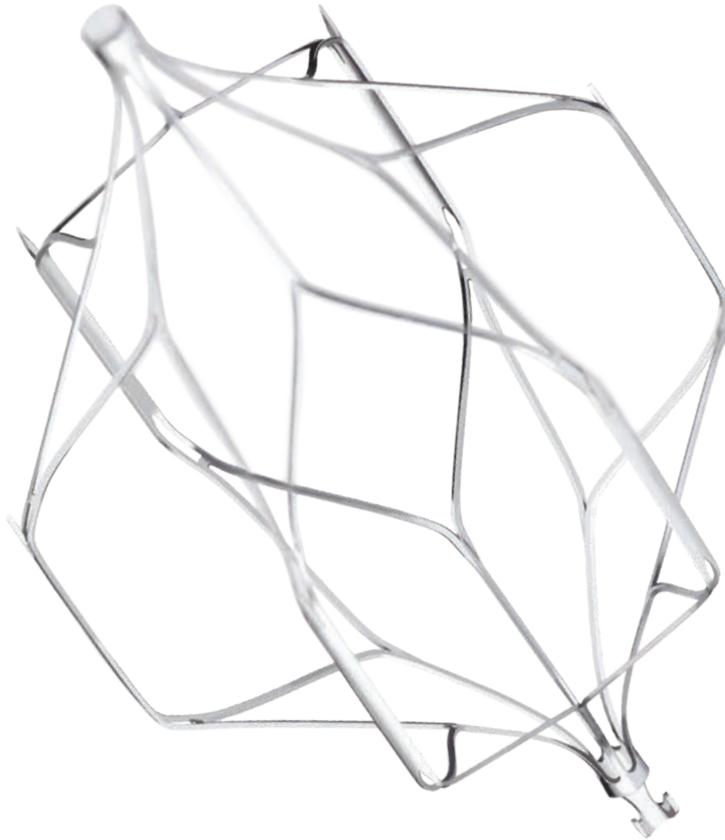


Side Struts
Self centering upon insertion minimizing the risk of tilting

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)	Must 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.



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

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.

For healthcare professionals only. Important information: Prior to use, refer to the instructions for use supplied with this device for indications, contraindications, suggested procedure, warnings and precautions. As part of its continuous product development policy, Cordis reserves the right to change product specifications without prior notification. Please contact your Cordis representative for additional product availability information. CORDIS, Cordis LOGO and OPTEASE are trademarks of Cordis and may be registered in the US and/or in other countries.
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