



PROXIMAL ANASTOMOSIS WITH
Enclose[®] II



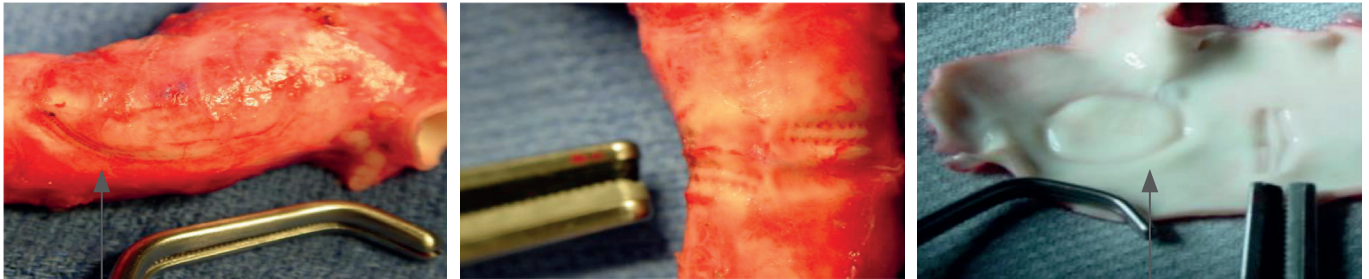
Go clampless

Proximal anastomosis assist device
for coronary bypass surgery

peters-surgical.com

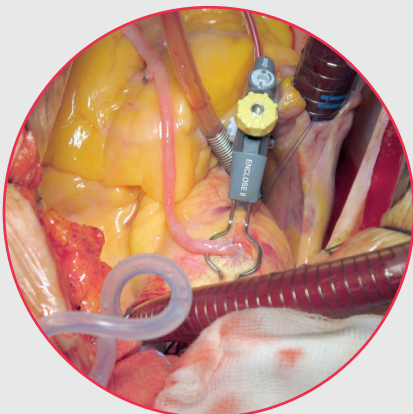
Enclose® II

Why is Clamping not preferred for proximal anastomosis with diffuse atherosclerotic plaque in ascending aorta ?



Aortic vessel trauma due to clamping

- There is a potential for plaque and tissue fragments to get embolized due to clamping, which can travel through the bloodstream and cause **cerebral embolization and stroke**.
- Embolization can occur in systemic circulation and can cause renal or mesenteric embolization causing **renal dysfunction or mesenteric ischemia**.
- **Aortic wall trauma or aortic wall dissection** can occur due to clamping.
- There can be an increased risk of post operative **cognitive dysfunction**.

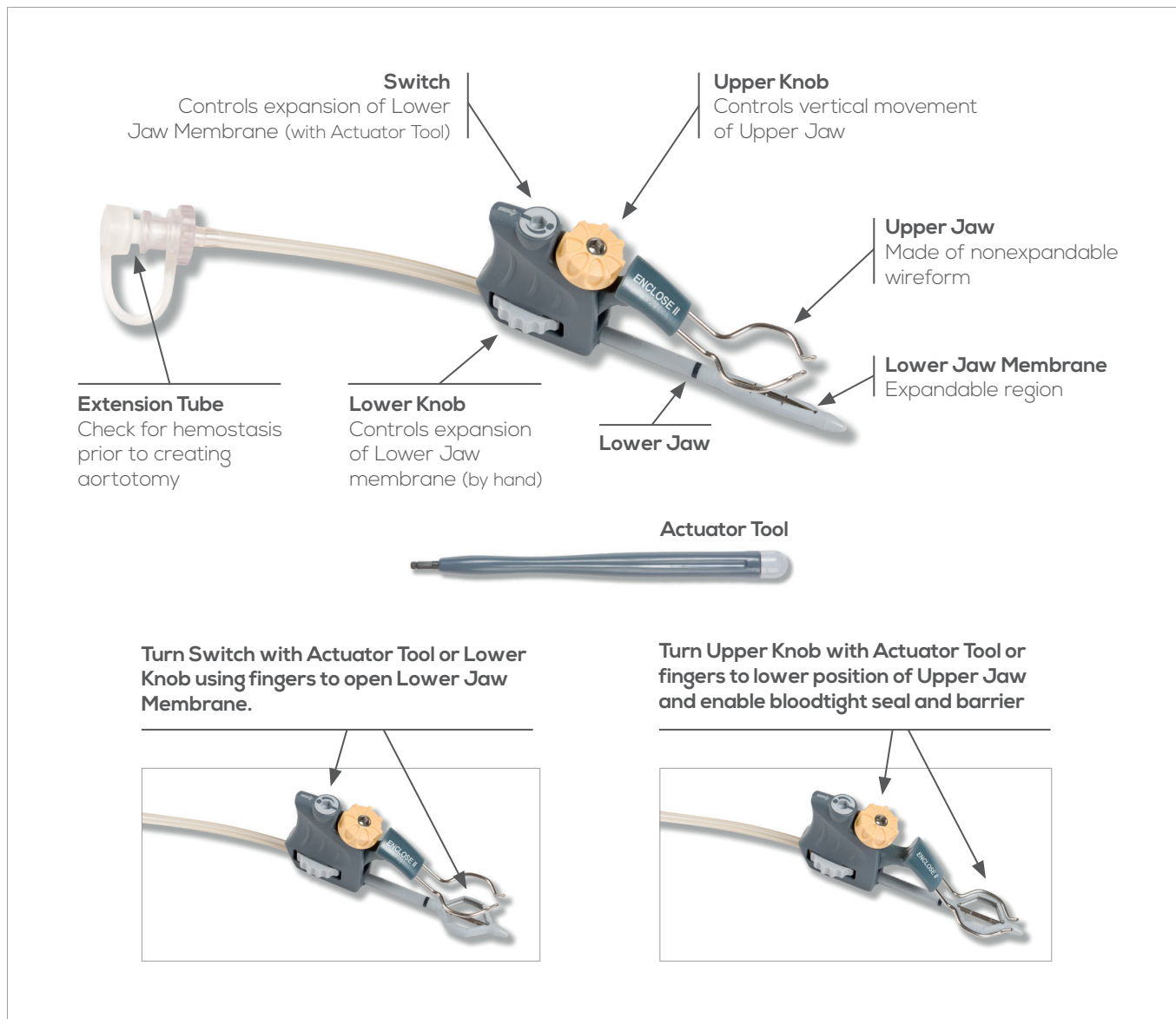


Enclose® II Anastomosis Assist Device

- Used in Beating Heart Coronary Bypass Surgery.
- Used in onpump CABG for those who perform proximal anastomosis after releasing the cross clamp.
- Designed specifically to obviate the need (and the associated risks) for partial clamping of the aorta.



Anatomy of the Enclose® II process



Confidence & Convenience for better outcomes

- Designed to provide the surgeon with a stable, bloodless field.
- Designed to eliminate the use of clamps, thus reducing cross-clamping and side clamping related complications.
- Manual anastomosis-assist device that allows hand sutured proximal anastomosis.
- Allows up several anastomosis from one insertion site.

Enclose® II



The Enclose® II Device

Item	Description	Aortic punch size	Contents
EN235V	Enclose Kit	3.5 mm	<ul style="list-style-type: none"> • Enclose Device • Actuator Tool • 14G Needle
EN240V	Enclose Kit	4.0 mm	<ul style="list-style-type: none"> • Enclose Device • Actuator Tool • 14G Needle
EN245V	Enclose Kit	4.5 mm	<ul style="list-style-type: none"> • Enclose Device • Actuator Tool • 14G Needle

ACTUATOR TOOL



NEEDLE



AORTIC PUNCH



The Enclose® II is included in the Peters Surgical product lines

INDICATIONS

The Enclose® II device is intended for use by cardiac surgeons during on-pump or off-pump coronary artery bypass grafting (CABG) procedures in place of partial occlusion clamps in ascending aortas free of atheromatous disease.

Manufacturer: Vitalitec International Inc d/b/a Peters Surgical USA. Distributed by PETERS SURGICAL

Presentation dedicated to PETERS SURGICAL employees, distributors and healthcare professionals.

Carefully read the Instructions For Use of medical device before use.



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