Instructions for Use

Reusable Attachments
for Universal Stabilizer Arms
Catalog Numbers 400-441, 400-442, 400-443, 400-451, 400-452, 400-453

Non-Sterile

Product Description

Universal Stabilizer Arm Accessory Attachments are reusable accessory devices for use with the ESTECH 401-161 Universal Stabilizer Arms™. The Attachments provide cardiac retraction and surgical exposure.

Use of Device

1. Attach Accessory Device to the ESTECH 401-161 Universal Stabilizer Arms™ by sliding sleeve of the quick connection away from flexible arm and inserting Attachment Shaft into the connector with the detent lined up to the screw hole. See Figures 1 and 2.

2. Position Device as required and as per instructions for use of the ESTECH 401-161 Universal Stabilizer Arms™.

CLEANING

Disassemble and scrub the instruments and parts thoroughly using a soft brush and mild detergent. Remove all traces of blood and debris. Make sure all parts are cleaned thoroughly. It is recommended that the instruments and parts be ultrasonically cleaned.

STERILIZATION

The instruments may be steam or gas sterilized. Refer to the sterilizer manufacturer's instructions for correct time, temperature and pressure settings.

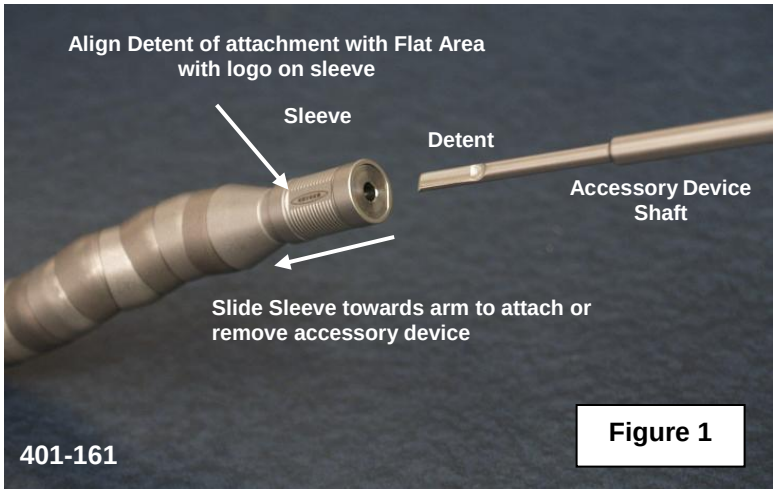

Figure 1

WARRANTY

AtriCure warrants that reasonable care has been used in the design and manufacture of this instrument. This warranty is in lieu of and excludes all other warranties not expressly set forth herein, whether expressed or implied by operation of law or otherwise, including, but not limited to, any implied warranties of merchantability or fitness for a particular use. Handling, storage, cleaning and

sterilization of this instrument as well as other factors relating to the patient, diagnosis, treatment, surgical procedures, and other matters beyond AtriCure control directly affect the instrument and the result obtained from its use. AtriCure's obligation under this warranty is limited to the repair or replacement of this instrument and AtriCure shall not be liable for any incidental or consequential loss, damage, or expense directly or indirectly arising from the use of this instrument. AtriCure neither assumes, nor authorizes any other person to assume for it, any other or additional liability or responsibility in connection with this instrument. AtriCure assumes no liability with respect to instruments reused, reprocessed or re-sterilized and makes no warranties expressed or implied, including but not limited to merchantability or fitness for intended use, with respect to such instrument.

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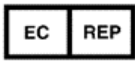

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Graphic Symbols for Device Labeling

Catalogue Number 	Batch Code 	Quantity 	Date of Manufacture 
Manufacturer/Company Address 	Authorized Representative in the European Union 	Instrument supplied Non-Sterile 	
Does not contain Natural Rubber Latex 	Contains no di (2-ethylhexyl) phthalate (**DEHP**) released from polyvinyl chloride (PVC) 	Consult Instructions For Use 	

CE mark, Product conforms with the essential requirements in the European Medical Devices Directive 93/42/EEC. CE	Caution: U.S. Federal law restricts this device to sale by or on the order of a physician or other licensed practitioner. ℞ ONLY