Quattro[®] X Suture Anchor with BroadBand[™] Tape

for Medial Row Fixation in Rotator Cuff Repair

Surgical Technique





Quattro X Suture Anchor with BroadBand Tape

Quattro X Suture Anchors are made of PEEK (PolyEther-EtherKetone) and available in 5.5 mm and 6.5 mm sizes.

- "Non-sliding" and "Sliding" BroadBand Tape options are also available.
- "Non-sliding" Quattro X Anchors (CM-9255BG and CM-9265BG) are double-loaded with one (1) non-sliding (blue) and one (1) sliding (white/green) BroadBand Tape. The "Sliding" Quattro X Anchors (CM-9255BGS and CM-9265BGS) are double-loaded with two (2) sliding BroadBand Tape strands (blue and white/ green).

Potential "Non-Sliding" Anchor Techniques:

- Double Row Knotless Repairs
- Medial Pulley Technique
- Medial Rip Stop Technique

Potential "Sliding" Anchor Techniques:

- Single Row Repairs
- Double Row Knotted Repairs

Instructions for Use

Quattro X Anchor instruments should be used to ensure the proper insertion of the Quattro X Anchor.

Quattro X Suture Anchor Instruments

Soft Bone	5.5/6.5mm Quattro X Awl (CM-9200), Purple Handle	
Hard Bone	5.5mm Quattro X Tap (CM-9201), Purple Handle	
	6.5mm Quattro X Tap (CM-9202), Purple Handle	



Site Preparation

Step 1

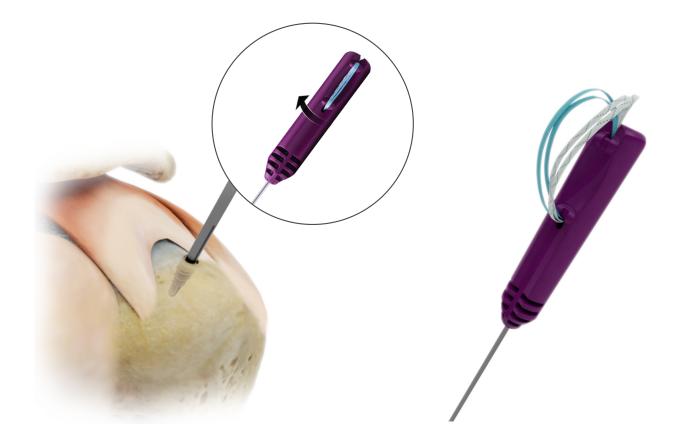
Use a shaver/burr/rasp to remove soft tissue directly overlying the insertion site.

Create Pilot Hole

Step 2

Place the Quattro X Anchor Awl (5.5/6.5mm) perpendicular to the bone at the medial edge of the anatomic footprint. Mallet the awl until the first horizontal laser-etch line is flush with the bone for the 5.5mm anchor and the second horizontal laser-etch line for the 6.5mm anchor.

Note: For hard bone, use the appropriate Quattro X Anchor Tap (5.5mm or 6.5mm) to create the pilot hole.



Insert Quattro X Anchor

Step 3

Insert the tip of the Quattro X Anchor into the prepared hole. Screw the Quattro X Anchor clockwise until the horizontal laser-etch line on the inserter shaft is flush with the bone surface.

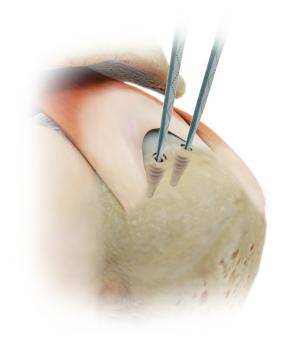
Note: For proper suture orientation, confirm that the vertical laser-etch line on the inserter shaft is facing or adjacent to the edge of the tissue.

Release Sutures

Step 4

Pull the tape strands completely out of the inserter handle and release from cleat.





Remove Inserter

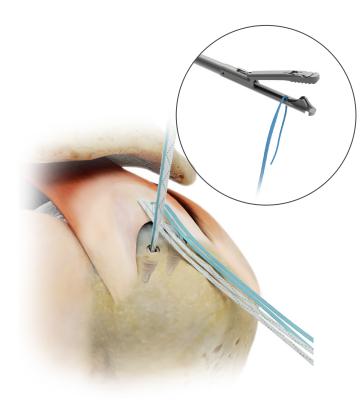
Step 5

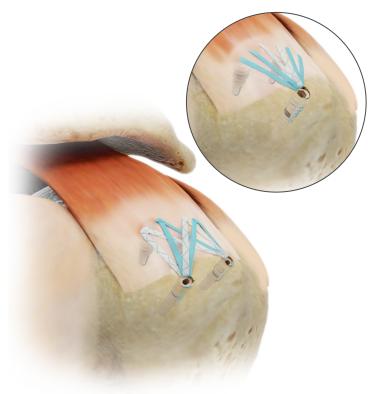
Pull back slowly on the inserter handle to disengage the inserter shaft from the anchor.

Insert Second Medial Row Anchor (Optional)

Step 6

Repeat steps 1-5 to insert a second medial row Quattro X Anchor (if desired).





Pass Suture

Step 7

Pass the BroadBand Suture Tape limbs through the tissue using a Quattro GT Suture Passer (CM-9010GT).

Finalize Repair

Step 8

Complete the repair with the preferred technique.

For Arthroscopic Knot Tying:

Always tie the blue tape strand first when using the "sliding" anchors to allow both BroadBand tape strands to slide easily.

For Double Row Repairs:

Load up to six (6) BroadBand Tape limbs into a Ventix[™] Link or Quattro Link Knotless Anchor to finalize the repair (refer to the Ventix Link and Quattro Link Anchor Surgical Techniques for complete implantation surgical instructions)

Ordering Information

Quattro X Suture Anchors with BroadBand Tape

Part Number	r Description	
CM-9255BG 5.5mm Quattro X Suture Anchor Double-Loaded with BroadBand Tape (Non-SI		
CM-9265BG	6.5mm Quattro X Suture Anchor Double-Loaded with BroadBand Tape (Non-Sliding)	
CM-9255BGS	5.5mm Quattro X Suture Anchor Double-Loaded with BroadBand Tape (Sliding)	
CM-9265BGS	6.5mm Quattro X Suture Anchor Double-Loaded with BroadBand Tape (Sliding)	

Quattro X Suture Anchor Reusable Instruments (Non-Sterile)

Part Number	Description
CM-9200	Awl, 5.5/6.5mm Quattro X Suture Anchor
CM-9201	Tap, 5.5mm Quattro X Suture Anchor
CM-9202	Tap, 6.5mm Quattro X Suture Anchor

Indications & Contraindications

INDICATIONS

The Quattro X Suture Anchors with BroadBand Tape are intended to be used for the reattachment of soft tissue to bone in rotator cuff repair.

CONTRAINDICATIONS

- 1) Surgical procedures other than those listed in the INDICATIONS section.
- 2) Presence of infection.
- 3) Patient conditions including insufficient or immature bone.
- 4) Insufficient blood supply or previous infections which may hinder the healing process.
- 5) Foreign body sensitivity. If material sensitivity is suspected, testing should be completed prior to device implantation.
- 6) The physician should carefully assess the device within immature bone.
- The placement of this device should not impact or disrupt the growth plate.
- 8) Conditions which may limit the patient's ability or willingness to follow postoperative care instructions.

All content herein is protected by copyright, trademarks and other intellectual property rights, as applicable, owned by or licensed to Zimmer Biomet or its affiliates un less ot herwise in dicated, and must not be redistributed, duplicated or disclosed, in whole or in part, without the express written consent of Zimmer Biomet. This material is intended for health care professionals. Distribution to any other recipient is prohibited. For indications, contraindications, warnings, precautions, potential adverse effects a n d p a tient c o unseling i n formation, s e e the package insert or contact your local representative; visit www. zimmerbiomet.com for additional product information.

Zimmer Biomet does not practice medicine. This technique was developed in conjunction with health care professionals. This document is intended for surgeons and is not intended for laypersons. Animations and virtual reality are provided as a visual guide based on surgical techniques; a written copy of the surgical technique is available at www. zimmerbiomet.com or from your local representative. Each surgeon should exercise his or her own independent judgment in the diagnosis and treatment of an individual patient, and this information does not purport to replace the comprehensive training surgeons have received. As with all surgical procedures, the technique used in each case will depend on the surgeon's medical judgment as the best treatment for each patient. Results will vary based on health, weight, activity and other variables. Not all patients are candidates for this product and/or procedure. Caution: Federal (USA) law restricts this device to sale by or on the order of a surgeon. Rx only.

©2021 Zimmer Biomet.

ECREP Authorized Representative

BIOMET GSCC B.V. Hazeldonk 6530 4836 LD Breda The Netherland



Legal Manufacturer Cayenne Medical Inc. 16597 N. 92nd St, Suite 101 Scottsdale, AZ 85260 USA

CE 2797

CE mark on a surgical technique is not valid unless there is a CE mark on the product label.

0603.1-US-en-REV1218

www.zimmerbiomet.com