Clinical Summary

Chronic Midsubstance Patellar Tendon and Retinacular Rupture: Primary Repair Enhancement Using Bioinductive Implant Augmentation

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Aim:

To describe the surgical technique and indications for using BioBrace® biocomposite scaffold in the repair of chronic midsubstance patellar tendon and retinacular ruptures.

Indications for Patellar Tendon Repair with Biobrace®

- Chronic Midsubstance Patellar Tendon Ruptures: Where tendon-to-tendon healing is required.
- Poor Tissue Quality: Cases with midsubstance tears and poor host tissue.
- Retinacular Tears: Medial and lateral retinacular injuries requiring augmentation.

Surgical Techniques

- Patient Positioning: Supine with a bump placed under the ipsilateral hip away from the sciatic nerve and a bone foam places under the injured knee.
- Incision and Dissection: Midline incision, centered over the patella and extending distally to the tibial tubercle.
- Tendon Fixation: Horizontal mattress sutures through tendon remnants and 23x30mm BioBrace[®], ensuring proper alignment and tension.
- Retinacular Repair: A 5x250 BioBrace[®] is placed in the retinacular injury both medically and laterally. It is trimmed to fit the appropriate distance and then sewn in with a #2 Dynacord.

Postoperative Rehabilitation

- Post Op Management: 6 weeks in a hinged knee brace with the knee locked in extension to help facilitate tendon healing.
- Gradual Mobilization: Followed by a structured rehabilitation protocol to restore function and gradual weight bearing.

Key Takeaways

- Enhanced Healing: BioBrace® promotes rapid cellular infiltration and tissue growth.
- Mechanical Support: Provides load-sharing strength at time zero, reducing re-rupture risk.
- Improved Outcomes: Early results show promising restoration of tendon strength.
- Future Potential: BioBrace® may be widely applicable for various tendon and ligament repairs.

