

GMReis Expands Its FDA-Cleared Trauma Portfolio with the 3.5 mm VersaLock® Tibial Plateau Plating System

July 23, 2026 | GMReis is proud to announce that the VersaLock® 3.5 mm Tibial Plateau Plating System has received FDA 510(k) clearance (K261406), marking another important milestone in the company's continued expansion in the U.S. orthopedic market.

This latest clearance further strengthens the VersaLock® Trauma Platform, providing surgeons with anatomically designed titanium implants featuring $\pm 15^\circ$ variable-angle locking technology and 3.5 mm dynamic compression for the treatment of tibial plateau fractures, osteotomies, and nonunions.

With this approval, GMReis now offers one of the most comprehensive FDA-cleared trauma plating portfolios available from a single manufacturer, covering virtually every major anatomical region, including:

- Foot & Ankle
- Tibial Plateau
- Proximal & Distal Femur
- Hand & Wrist
- Forearm
- Elbow
- Distal & Proximal Humerus
- Clavicle
- Scapula

The VersaLock® platform reflects GMReis' commitment to delivering innovative fixation technologies that combine anatomical design, advanced locking performance, and procedural versatility to meet the evolving needs of orthopedic trauma surgeons worldwide.

As GMReis continues to expand its U.S. portfolio, the company remains focused on providing comprehensive orthopedic solutions backed by manufacturing excellence, rigorous quality standards, and continuous innovation.

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