VIKING MIS PEDICLE SCREW SYSTEM

PRESIDIO SURGICAL

SLENDER PERCUTANEOUS EXTENSION FOR MINIMAL INCISION SIZE

SINGLE STEP EXTENSION REMOVAL

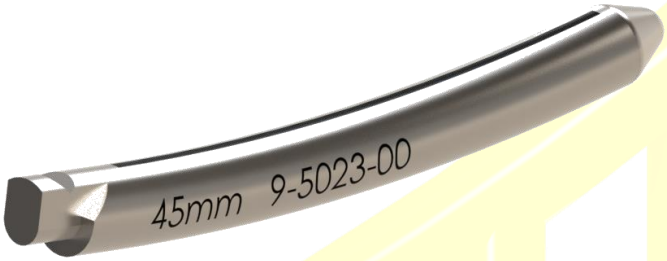
LOW PROFILE TULIP

FRICTION FIT TULIP FOR EASY POSITIONING AND ROD CAPTURE

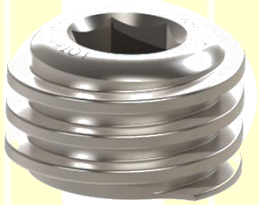
PROXIMALLY TAPERED CORE FOR INCREASED PURCHASE

SIZES:

5.0mm
6.0mm
7.0mm
8.0mm



SECURE ROD HOLDER FOR SAFE ROD PLACEMENT



SINGLE PIECE LOCKING SET SCREW FOR INCREASED HOLDING STRENGTH

RODS	
ITEM #	DESCRIPTION
9-5023-01	40mm Rod, Curved
9-5023-02	45mm Rod, Curved
9-5023-03	50mm Rod, Curved
9-5023-04	55mm Rod, Curved
9-5023-05	60mm Rod, Curved
9-5023-06	70mm Rod, Curved
9-5023-07	80mm Rod, Curved
9-5023-08	90mm Rod, Curved
9-5023-09	100mm Rod, Curved
*Larger sizes upon request	

SCREWS	
ITEM #	DESCRIPTION
9-5017-02	5.0 X 35mm Screw
9-5017-03	5.0 X 40mm Screw
9-5017-04	5.0 X 45mm Screw
9-5017-08	6.0 X 35mm Screw
9-5017-09	6.0 X 40mm Screw
9-5017-10	6.0 X 45mm Screw
9-5017-11	6.0 X 50mm Screw
9-5017-12	6.0 X 55mm Screw
9-5017-21	7.0 X 35mm Screw
9-5017-22	7.0 X 40mm Screw
9-5017-23	7.0 X 45mm Screw
9-5017-24	7.0 X 50mm Screw
9-5017-25	7.0 X 55mm Screw
9-5017-33	8.0 X 35mm Screw
9-5017-34	8.0 X 40mm Screw
9-5017-35	8.0 X 45mm Screw
9-5017-36	8.0 X 50mm Screw

PRESIDIO
SURGICAL

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Indications: The Viking Pedicle screw system is intended to provide immobilization and stabilization of spinal segments in skeletally mature patients as an adjunct to fusion in the treatment of the following acute and chronic instabilities or deformities of thoracic, lumbar, and sacral spine: degenerative disc disease(DDD) (defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies.); spondylolisthesis; trauma (i.e., fracture or dislocation); spinal stenosis; tumor; pseudoarthosis; and failed previous fusion.

Products Patented and Patents Pending

All products are not currently available in all markets

Contraindications: Disease conditions that have been shown to be safely and predictably managed without the use of internal fixation devices are relative contraindications to the use of these devices. Active systemic infection or infection localized to the site of the proposed implantation are contraindications to implantation. Severe osteoporosis is a relative contraindication because it may prevent adequate fixation of spinal anchors and thus preclude the use of this or any other spinal instrument system. Any entity or conditions that totally precludes the possibility of fusion, i.e., cancer, kidney dialysis, or osteopenia is a relative contraindication. Other relative contraindications include obesity, certain degenerative diseases, and foreign body sensitivity. In addition, the patient's occupation or activity level or mental capacity may be a relative contraindication to this surgery. Specifically, patients who because of their occupation or lifestyle, or because of conditions such as mental illness, alcoholism, or drug abuse, may place undue stresses on the implant during bony healing and may be at higher risk for implant failure.