

Aleutian[®]

Lateral Interbody System



The Aleutian Lateral Interbody System is designed to work in concert with the Ravine or Niagara Lateral Access Systems. Aleutian Lateral provides anterior column support, bridging of the disc space, and includes a full line of instrumentation designed specifically for the far lateral transpoas approach. Aleutian implants are intended to be used with supplemental internal fixation.

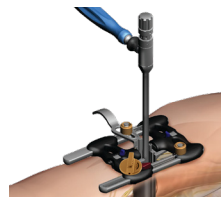
Aleutian Lateral Interbody System

The Interbody with a leading edge

- 6 mm bulletted nose allows for ease of insertion into collapsed disc spaces
- Circumferential rings designed to grip the endplates in all directions
- Large center channel designed for spacious bone graft
- Manufactured of biocompatible PEEK

Wide range of footprints available

- Widths available in 18mm and 22mm
- Available in 0°, 8°, 12° and 15° lordotic sagittal profiles
- Heights from 8 - 16mm and lengths from 45 - 60mm
- Add-on set with 16mm wide implants for smaller disc space



Multiple access options:

- Ravine, a dual flat blade, spine-based retractor directly fixated to the spine
- Niagara, a table-mounted tubular retractor allowing for control through precision, flexibility and visibility

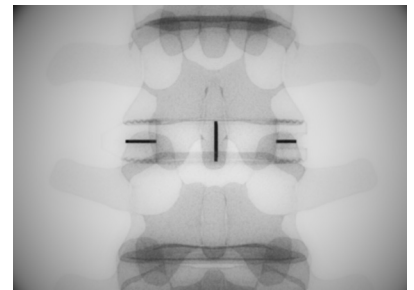
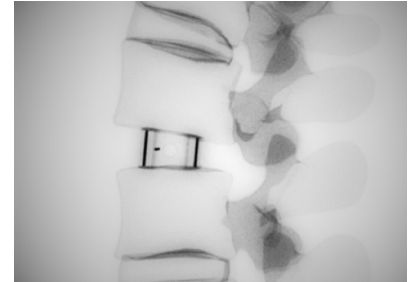


Instrument design features

- All instrumentation designed with working lengths for a lateral procedure, while facilitating a less obstructed view of the working channel
- Lateral Interbody Inserter with a rotatable offset handle allows for enhanced control
- Multiple graft containment ramps options available: integrated to the inserter or loose



Tantalum markers for placement confirmation



Dedicated versatile disc preparation instruments



Spine division

This document is intended solely for the use of healthcare professionals. A surgeon must always rely on his or her own professional clinical judgment when deciding whether to use a particular product when treating a particular patient. Stryker does not dispense medical advice and recommends that surgeons be trained in the use of any particular product before using it in surgery.

The information presented is intended to demonstrate the breadth of Stryker product offerings. A surgeon must always refer to the package insert, product label and/or instructions for use before using any Stryker product. The products depicted are CE marked according to the Medical Device Directive 93/42/EEC. Products may not be available in all markets because product availability is subject to the regulatory and/or medical practices in individual markets. Please contact your Stryker representative if you have questions about the availability of Stryker products in your area.

Stryker Corporation or its divisions or other corporate affiliated entities own, use or have applied for the following trademarks or service marks: Aleutian, K2M, Ravine, Serengeti, Stryker. All other trademarks are trademarks of their respective owners or holders.

ALELA-SS-1_25381
Copyright © 2020 Stryker



K2M, Inc.
600 Hope Parkway SE
Leesburg, VA 20175
t: 571 919 2000

www.stryker.com