





International

The MectaLIF System is indicated for use in skeletally mature patients suffering from Degenerative Disc Disease at one level or two contiguous levels from L2 to S1. Patients should have received 6 months of non-operative treatment prior to treatment with the devices.

MECTALIF SYSTEM

ANATOMICAL DESIGN

- Shaped to facilitate proper load transfer and enhance the contact surface
- Bioconvex surface to match endplate anatomy
- Bi-cortical bridge to offer a stable support and reduce the risk of subsidence

STABILITY FEATURES

• Pyramid teeth are designed to grip the endplate and to help limit implant migration

• Broad surface is designed to contribute to mechanical stability under lateral bending and to provide anterior / posterior rim bridging



TIPEEK TECHNOLOGY

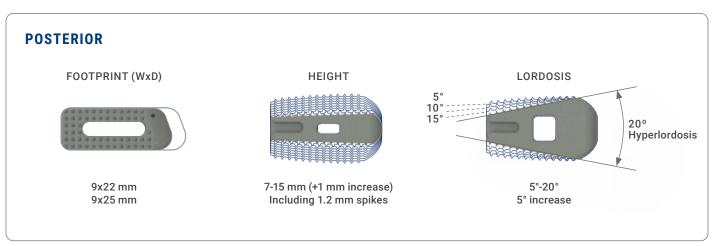
The MectaLIF system includes plasma-sprayed titanium coated cages.

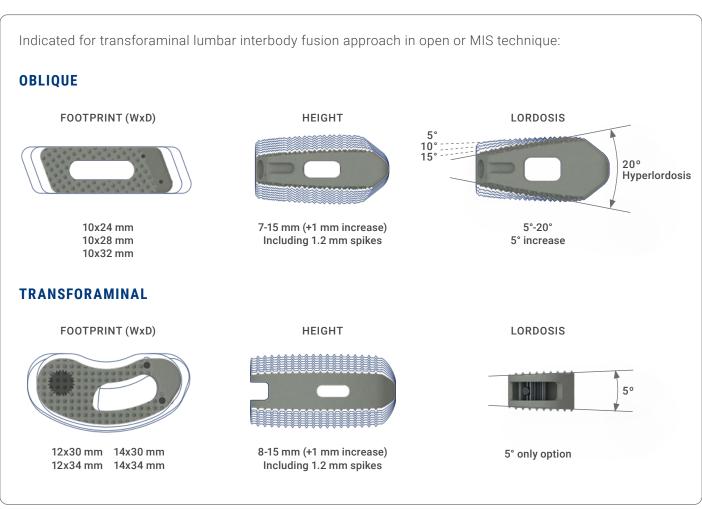
The TiPEEK osteoconductivity in the MectaLIF Cages enhances primary stability due to complex micro rough surface that improves friction increasing migration resistance.^[1,2]



A COMPREHENSIVE SYSTEM

The MectaLIF family of interbody fusion devices offers a variety of solutions to cover patient's unique anatomies and surgical needs:







AectaLIF® SYSTEM

EASE OF IMPLANTATION

All cages included in the MectaLIF system are equipped with a self-distracting bullet nose.

Dedicated instruments for each cage family simplify the surgical steps:







References: [1] M.Rickert et al. Transforaminal lumbar interbody fusion in PEEK oblique cages with and without titanium coating: results from a randomized clinical trial 8th M.O.R.E. International Symposium. [2] B.Walsh et al. Titanium coated interbody devices 8th M.O.R.E. International Symposium.

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