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NEWS RELEASE FOR IMMEDIATE RELEASE

Theken Announces First Surgery Performed in Baltimore for New Spinal Fusion System

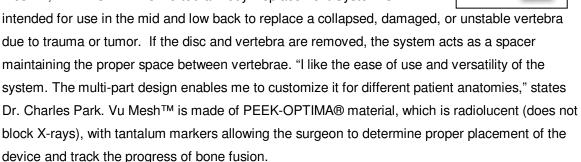
Vu Mesh™ PEEK-OPTIMA Vertebral Body Replacement receives FDA Clearance

Akron, OH – (May 16, 2007) Theken Spine, a leader in spinal fixation systems, announces that the first surgery for the Vu Mesh system was performed by Charles Park, MD in a leading hospital in Baltimore, MD.

Theken received FDA clearance to market Vu Mesh in the United States in late April. This surgery kicks off the alpha phase. This is the first product out of six for which Theken anticipates receiving FDA clearance this year.

The Vu Mesh product launch is targeted for mid-Summer 2007.

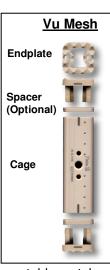
Theken worked with a team of design surgeons to develop this product. Vu Mesh $^{\text{TM}}$, PEEK-OPTIMA® Vertebral Body Replacement System is



For up to date information, contact Rachael Smith rsmith@thekenspine.com or 330-807-2569.

About Theken, a family of companies

The Theken family of companies Theken Spine, LLC, Theken Disc, LLC, Theken Orthopaedic, Inc. and Therics, LLC (www.theken.com) specializes in pioneering spinal implant technologies that improve spinal surgical techniques benefiting patients as well as surgeons. Theken provides comprehensive product lines that offer surgeons peace of mind through steadfast product reliability and easy-to-use instrumentation. Products include cervical plates, pedicle screws,



spacers, degenerative/deformity and trauma devices. Theken also leads the market in next-generation artificial disc replacement technology.

Company Contact

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