

Manufactured by:
Southern Medical (Pty) Ltd
Building 10,
Southern Implants Office Park,
1 Albert Road,
Irene, 0062
South Africa
Tel: +27 12 667 6243/4
Email: info@southmed.co.za

European Representative:
Southern Implants UK, Inc.
Building 3, Chiswick Park

566 Chiswick, High Road
London W4 5YA, United Kingdom

uncertain be sure to contact a Southern Medical Representative.

Surgical Risks:

- Abdominal hernia
- Allergic or other reaction to anesthesia
- Annular ossification
- Blood loss or hemorrhage
- Death
- Degenerative changes in adjacent segment
- Dural injury
- Facet joint deterioration
- Hematoma or seroma
- Heterotopic ossification
- Ileus
- Implant breakage
- Implant collapse or subsidence into adjacent vertebrae
- Implant degradation
- Implant displacement/migration
- Impotence
- Incontinence
- Infection
- Kidney or ureter injury
- Metal ion release
- Myocardial infarction
- Nerve root injury
- Neurologic deterioration:
 - Clumsiness
 - Foot drop
 - Limp
 - Short step
 - Slow moving gait
 - Weakness
- Numbness

- Osteolysis or vertebral inflammation
- Osteophyte resorption
- Pain
- Perineural fibrosis
- Peritonitis
- Pneumonia
- Pneumothorax
- Pulmonary embolism
- Removal of the device in the post-op or follow-up period
- Reoperation at the study treatment level with or without removal or modification of any or all components of the device
- Retrograde ejaculation
- Revision with or without replacement of a component
- RSD (reflex sympathetic dystrophy)
- Spinal instability
- Spinal stenosis (narrowing of the spinal canal)
- Spondylolisthesis acquisita
- Spondylosis acquisita
- Spontaneous fusion
- Sterility
- Supplemental fixation
- Surgical instrument failure
- Thrombosis
- Tumor formation/ carcinogenesis potential
- Vertebral fracture
- Vessel damage
- Wear debris generation

RE-STERILIZATION AND RE-USE WARNING:



Do not re-sterilize implants produced sterile. Implants are sterilized by gamma irradiation. Re-sterilization could cause material degradation and could result in surgical rejection and/or post-operative infection. The implant is designed for single patient use only and must never be reused. An explanted implant must never be re-implanted. Reuse or re-implantation may result in cross-contamination or infection.

Magnetic Resonance Imaging (MRI)

The TLIF devices have not been tested for adverse effects under MRI. The TLIF is manufactured from non-ferromagnetic materials. Non-clinical evaluations have demonstrated that TLIF devices are MR Conditional based on a Cobalt Chrome worst case device. Patients can be scanned safely immediately after implantation under the following conditions:

- Static magnetic field of 3.0-Tesla (3.0T) or less
- Maximum spatial gradient field less than or equal to 10T/m.
- Normal Operating Mode: Maximum whole-body specific absorption rate (SAR) of:
 - 2 W/kg for 15 minutes of scanning at 1.5T.
 - 2 W/kg for 15 minutes of scanning at 3.0T.












MR image quality may be compromised if the area of interest is the same or relatively close to the position of the device, and it may be necessary to optimize the MR imaging parameters.

Risks of placing implants in or near a magnetic field include: (1) movement of ferromagnetic components, (2) localised heating of components caused by radio frequency induction heating and (3) image artifacts created by interaction between metallic components and the magnetic field.

Post Implantation:

The surgeon/physician's postoperative directions, indications and warnings to the patient with subsequent patient compliance are vital to successful fusion. The subject must refrain from physical activity for several months. Physical movement must be minimized. Subjects will be instructed not to engage in activities requiring lifting, bending, twisting or excessive movement for several months post-operatively. The subject must not be exposed to electrical shock and/or mechanical vibrations. Directly after the operation, subjects will be placed in a brace until fusion occurs. After a couple of months, the subject may return to the workplace if the work environment does not entail excessive physical activities. Car journey duration must be minimized, preferably avoided.

Descriptions of Symbols Used in Packaging:

USE BY	
LOT NUMBER	
DATE OF MANUFACTURE	
MANUFACTURER ADDRESS	
DO NOT REUSE	
STERILIZED USING IRRADIATION	
CAUTION	
AUTHORISED REPRESENTATIVE IN THE EUROPEAN COMMUNITY	
CONSULT THE INSTRUCTIONS FOR USE	 southmed.co.za
DO NOT RESTERILIZE	
DO NOT USE IF PACKAGING IS DAMAGED	

IMPORTANT: PLEASE READ

For detailed information on the Southern Transforaminal Interbody Fusion (TLIF), please consult the Surgeons' Manual and IFU at southmed.co.za

Description:

The Bullet Transforaminal Lumbar Interbody Fusion (TLIF) is a single component device, manufactured from biocompatible poly-ether-ether-ketone (PEEK) (ASTM F2026) and tantalum (ASTM F560). The device serves to augment posterior pedicle screw fixation. Surgical instrumentation is manufactured from surgical grade stainless steel (ASTM F899).

The Camber Transforaminal Lumbar Interbody Fusion (TLIF) is manufactured from biocompatible poly-ether-ether-ketone (PEEK) (ASTM F2026) with tantalum (ASTM F560) markers and a titanium (ASTM F136) swivel joint. The device serves to augment posterior pedicle screw fixation. Surgical instrumentation is manufactured from surgical grade stainless steel (ASTM F899).

The Curved Transforaminal Lumbar Interbody Fusion (TLIF) is manufactured from biocompatible poly-ether-ether-ketone (PEEK) (ASTM F2026) with tantalum (ASTM F560) markers. The device serves to augment posterior pedicle screw fixation. Surgical instrumentation is manufactured from surgical grade stainless steel (ASTM F899).

Variants: All TLIF cages are available in an additional configuration whereby the vertebral contacting surfaces are titanium (ASTM F1580) plasma coated.

Radioactivity warning:

No radioactive substance or radioactivity.

Intended purpose:

The intervertebral endoprosthesis intended as a treatment option for pain and functional disorders specific to the lumbar vertebral column. The aim of the device is to provide additional support for pedicle screw fixation between two vertebral bodies and initial immobilization of these bodies whilst simultaneously providing space for bone graft so that a fusion of the two vertebral bodies will in time be attained.

Intended performance and undesirable side-effects:

Preferred patients are those that have instability due to degenerated discs and/or facet joints causing pain, loss of disc height, spondylolisthesis or change in the normal curvature of the spine. The Bullet TLIF is intended for use with bilateral pedicle screw fixation in the lumbar spine, including L5-S1. Revision surgery for device retrieval or additional instrumentation must be possible in the event of failure to fuse or poor clinical outcome.

Indications:

- Age between 18 and 60 years
- Single level degenerative disc disease and instabilities
- Fractures
- Mechanical back pain (usually due to disc degeneration)
- Pseudarthrosis or failed arthrodesis
- Recurrent disc herniation
- Scoliosis
- Segment instability (bulging disc)
- Some tumors
- Spinal instability (spondylolisthesis)
- Spinal Stenosis

Contraindications:

- Certain spinal fractures
- Certain spinal tumors
- Major mental illnesses and psychosocial disorders (Waddell>3/5)
- Primary spinal deformity
- Serious instabilities
- Spinal fractures
- Treatment at L5-S1 level (for Curved and Camber TLIF)

Sterility:

All implants are supplied sterile, and are for single use only before the labeled expiration date. Do not re-use implants. Do not use implants if the packaging has been damaged or previously opened, or if the expiration date has passed. If

Risks Associated with Lumbar Spinal Systems:

- Acute heart failure
- Annular ossification
- Degenerative changes in adjacent segment
- Dural injury
- Facet joint deterioration
- Hematoma or seroma
- Heterotopic ossification
- Implant breakage
- Implant collapse or subsidence into adjacent vertebrae
- Implant degradation
- Implant displacement
- Impotence
- Incontinence
- Kidney or ureter injury
- Metal ion release
- Nerve root injury
- Neurologic deterioration
- Clumsiness
- Foot drop
- Limp
- Short step
- Slow moving gait
- Weakness
- Numbness

- Osteophyte resorption
- Perineural fibrosis
- Removal of the device in the post-op or follow-up period
- Reoperation at the study treatment level with or without removal or modification of any or all components of the device
- Revision with or without replacement of a component
- Retrograde ejaculation
- RSD (reflex sympathetic dystrophy)
- Spinal cord injury due to instruments being forced too deep
- Spinal instability
- Spinal stenosis (narrowing of the spinal canal)
- Spondylolisthesis acquisita
- Spondylosis acquisita
- Spontaneous fusion
- Sterility
- Supplemental fixation
- Tumor formation/ carcinogenesis potential
- Vertebral fracture
- Vessel damage and
- Wear debris generation

Recommended Surgical Procedure:

Refer to the surgical procedure provided by Southern Medical (Pty) Ltd.

USAGE WARNING:

Improper technique in implant placement can result in implant failure. The TLIF devices are not intended as the sole means of spinal support. In absence of bone graft or fusion the implant or implant components can be expected to pull out, bend or fracture as a result of everyday mechanical stresses. Placement of devices is limited to surgeons. Refer to the surgical procedure and product brochure for more information.

