

Trust the
science.

InterSpace[®]
Knee Spacer



Demonstrating strong eradication rates before and after second-stage revisions.¹

19

Peer-reviewed
papers at 18
different sites

732

Cases
reported

94.1

Percent eradication
rate at implant
removal

93.8

Percent eradication
rate at latest
follow-up

55

Months average
follow-up

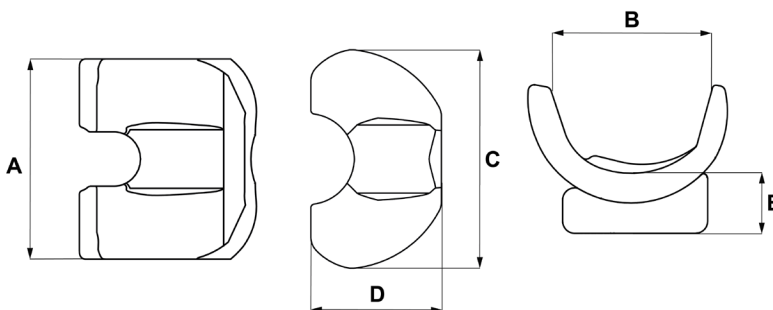
Surgeon focused. Patient driven.[™] **Exactech[®]**

InterSpace® Knee Spacer

InterSpace® Knee is designed to temporarily replace a Total Knee Arthroplasty (TKA) when the implant is removed as a result of infection. InterSpace Knee is a partial load-bearing structure consisting of Gentamicin-impregnated PMMA bone cement.

ADVANTAGES

- Proven high release formulation and design yields predictable and consistent local antimicrobial activity compared to other treatment options²⁻⁴
- Preformed spacers are designed to maintain joint space and allows limited mobility with partial weight bearing^{5-7*}
- Stabilizes or tensions the soft tissues and reduces bone loss between stages, potentially facilitating easier re-implantation during a second-stage procedure^{6,8-9}
- Improves quality of life between procedures^{6,8}
- Provides predictable, consistent local antibiotic release^{2,3}
- Preformed spacers have been shown to shorten operating room time¹⁰
- Reduces hospitalization and allows for an early transition to rehabilitation and physical therapy^{11,12}
- InterSpace demonstrates functional success rates equivalent to non-infected revisions.⁹



SIZE CHART

Size	REF	A (mm)	B (mm)	C (mm)	D (mm)	E (mm)	Gentamicin Base
Small (S)	SPK0022	54	40	60	36	16	0.9g
Medium (M)	SPK0122	64	47	70	42	17	1.3g
Large (L)	SPK0222	74	54	80	48	18	1.8g
Extra Large (XL)	SPK0322	84	61	90	54	19	2.7g

Trials

SPK90Z1 InterSpace Knee Trials (S, M, L)
SPK03Z1 InterSpace Knee Trials (XL)

*Partial weight bearing must be assessed on an individual basis with relation to the anatomic condition of the local bone, bone quality and clinical conditions of the patient during rehabilitation stages. Care must be taken to minimize the risk of damaging bone tissue and the implant through excessive weight bearing or forced mobilization.

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