

K073105 PEEK SPINAL SPHERESJan 3, 2008
62 days to decisionK073105 · Product code: **NVR** · Orthopedic
Source: <https://www.510kdatabase.net/k073105/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - U
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Solid-sphere, Lumbar (NVR)
Date received	Nov 2, 2007
Decision date	Jan 3, 2008
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interbody Innovations, Llp
Location	Clearwater, FL, US
Contact	J.D. WEBB
510(k) history	4 submissions · 1 cleared · 2005-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073105/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026