FDA 510(k) Premarket Notification

K051371 INTERBODY INNOVATIONS CEMENT RESTRICTOR

CLEARED

Dec 6, 2005
194 days to decision

K051371 · Product code: **JDK** · Orthopedic
Source: https://www.510kdatabase.net/k051371/

SUBMISSION DETAILS

Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	May 26, 2005
Decision date	Dec 6, 2005
Days to decision	194 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Interbody Innovations, Llp**
Location	Clearwater, FL, US
Contact	IAN GORDON
510(k) history	4 submissions · 1 cleared · 2005-2008