FDA 510(k) Premarket Notification

K072120 PARAMOUNT INTERVERTEBRAL BODY FUSION DEVICE

CLEARED

Oct 11, 2007
71 days to decision

K072120 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k072120/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 1, 2007
Decision date	Oct 11, 2007
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Innovative Spinal Technologies, Inc.**
Location	Mansfield, MA, US
Contact	GINA YEH
510(k) history	4 submissions · 4 cleared · 2005-2007