FDA 510(k) Premarket Notification

K080411 PLATEAU SPACER SYSTEM

CLEARED

May 15, 2008
90 days to decision

K080411 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k080411/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 15, 2008
Decision date	May 15, 2008
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Life Spine**
Location	Hoffman Estates, IL, US
Contact	REBECCA M BROOKS
510(k) history	36 submissions · 34 cleared · 2006-2018