FDA 510(k) Premarket Notification

K130573 TYBER MEDICAL INTERBODY SYSTEM

CLEARED

Sep 30, 2013
210 days to decision

K130573 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k130573/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Mar 4, 2013
Decision date	Sep 30, 2013
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary
Other names	ACIF, ALIF, PLIF, TLIF, DLIF

APPLICANT

Company	**Tyber Medical, LLC**
Location	Morristown, NJ, US
Contact	JEFF TYBER
Website	https://www.tybermed.com
510(k) history	28 submissions · 28 cleared · 2013-2026