FDA 510(k) Premarket Notification

K181261 Curiteva Cervical Interbody Fusion System

CLEARED

Jul 9, 2018
59 days to decision

K181261 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k181261/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	May 11, 2018
Decision date	Jul 9, 2018
Days to decision	59 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Curiteva, LLC**
Location	Tanner, AL, US
Contact	Eric Linder
510(k) history	4 submissions · 4 cleared · 2018-2018