FDA 510(k) Premarket Notification

K171444 YUKON OCT Spinal System

CLEARED

Aug 8, 2017
84 days to decision

K171444 · Product code: **NKG** · Orthopedic
Source: https://www.510kdatabase.net/k171444/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	May 16, 2017
Decision date	Aug 8, 2017
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**K2m, Inc.**
Location	Leesburg, VA, US
Contact	Nancy Giezen
510(k) history	100 submissions · 97 cleared · 2007-2026