FDA 510(k) Premarket Notification

K133924 ENDOLUMINAL OCCLUSION SYSTEM (EOS)

CLEARED

Dec 3, 2014
345 days to decision

K133924 · Product code: **KRD** · Cardiovascular
Source: https://www.510kdatabase.net/k133924/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Dec 23, 2013
Decision date	Dec 3, 2014
Days to decision	345 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Artventive Medical Group, Inc.**
Location	Woodinville, WA, US
Contact	Roberta Hines
510(k) history	3 submissions · 3 cleared · 2014-2022