FDA 510(k) Premarket Notification

K141141 KIVA VCF TREATMENT SYSTEM

CLEARED

Aug 14, 2014
104 days to decision

K141141 · Product code: **NDN** · Orthopedic
Source: https://www.510kdatabase.net/k141141/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Bone, Vertebroplasty (NDN)
Date received	May 2, 2014
Decision date	Aug 14, 2014
Days to decision	104 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Benvenue Medical, Inc.**
Location	Mountain View, CA, US
Contact	CINDY DOMECUS, R.A.C.
510(k) history	8 submissions · 8 cleared · 2007-2020