

**K141141 KIVA VCF TREATMENT SYSTEM**Aug 14, 2014  
104 days to decisionK141141 · 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	<b>Benvenue Medical, Inc.</b>
Location	Mountain View, CA, US
Contact	CINDY DOMECUS, R.A.C.
510(k) history	8 submissions · 8 cleared · 2007-2020

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141141/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026