

K082134 CIRCULATOR BOOTMay 7, 2009
282 days to decisionK082134 · Product code: **DRN** · Cardiovascular
Source: <https://www.510kdatabase.net/k082134/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Counter-pulsating, External (DRN)
Date received	Jul 29, 2008
Decision date	May 7, 2009
Days to decision	282 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Circulator Boot Corp.
Location	Mchenry, IL, US
Contact	RICHARD S DILLON
510(k) history	4 submissions · 4 cleared · 1980-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082134/> Data sourced from FDA 510(k) public records (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. - Generated June 23, 2026