FDA 510(k) Premarket Notification

K082134 CIRCULATOR BOOT

CLEARED

May 7, 2009
282 days to decision

K082134 · 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