FDA 510(k) Premarket Notification

K033657 MOBILE EXTERNAL COUNTER PULSATION SYSTEM ANGIONEW-V

CLEARED

Dec 19, 2003
28 days to decision

K033657 · Product code: **DRN** · Cardiovascular
Source: https://www.510kdatabase.net/k033657/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Counter-pulsating, External (DRN)
Date received	Nov 21, 2003
Decision date	Dec 19, 2003
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Living Data Technologies Corporation**
Location	Great Neck, NY, US
Contact	SUSAN D GOLDSTEIN-FALK
510(k) history	2 submissions · 2 cleared · 2003-2003