FDA 510(k) Premarket Notification

K792430 CARDIASSIST EXTERNAL COUNTERPULSATION

CLEARED

Jan 23, 1980
57 days to decision

K792430 · Product code: **DRN** · Cardiovascular
Source: https://www.510kdatabase.net/k792430/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Counter-pulsating, External (DRN)
Date received	Nov 27, 1979
Decision date	Jan 23, 1980
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Cardiassist Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980