FDA 510(k) Premarket Notification

K800179 INFRASONDE AUTO/INFLATOR ELEC. VIBRATOR

CLEARED

Feb 11, 1980
13 days to decision

K800179 · Product code: **DXW** · Cardiovascular
Source: https://www.510kdatabase.net/k800179/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Phonocatheter, Intracavitary (DXW)
Date received	Jan 29, 1980
Decision date	Feb 11, 1980
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Sphygmetrics, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1980