FDA 510(k) Premarket Notification

K800201 ESOPHAGEAL PILL-ELECTRODE

CLEARED

Feb 26, 1980
27 days to decision

K800201 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k800201/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 30, 1980
Decision date	Feb 26, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Arzbaecher & Co.**
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1979-1980