FDA 510(k) Premarket Notification

K852181 ELECTROCARDIOGRAPH

CLEARED

Oct 9, 1985
142 days to decision

K852181 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k852181/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	May 20, 1985
Decision date	Oct 9, 1985
Days to decision	142 days
Third-party review	No

APPLICANT

Company	**Armstrong Ind., Inc.**
Location	Mchenry, IL, US
Contact	KAREN BOWERS
510(k) history	5 submissions · 5 cleared · 1982-1985