FDA 510(k) Premarket Notification

K852281 SCHILLER EKG COMBI MSG-IM SCOPE & RECORDER

CLEARED

Oct 9, 1985
133 days to decision

K852281 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k852281/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	May 29, 1985
Decision date	Oct 9, 1985
Days to decision	133 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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k852281/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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