FDA 510(k) Premarket Notification

K840569 EKG 501 A

CLEARED

Dec 30, 1985
691 days to decision

K840569 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k840569/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Feb 8, 1984
Decision date	Dec 30, 1985
Days to decision	691 days
Third-party review	No

APPLICANT

Company	**Bosch Hearing Instruments, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1980-1985