

K840569 EKG 501 ADec 30, 1985
691 days to decisionK840569 · Product code: **DPS** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k840569/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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