

K760563 ECG MODULATORSep 10, 1976
8 days to decisionK760563 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k760563/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Sep 2, 1976
Decision date	Sep 10, 1976
Days to decision	8 days
Third-party review	No

APPLICANT

Company	Physio-Control Corp.
Location	Mchenry, IL, US
510(k) history	80 submissions · 78 cleared · 1976-1999

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Device record: <https://www.510kdatabase.net/k760563/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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