

K772350 PATIENT ALERT MODEL 555Mar 9, 1978
76 days to decisionK772350 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k772350/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Dec 23, 1977
Decision date	Mar 9, 1978
Days to decision	76 days
Third-party review	No

APPLICANT

Company	Intermedics, Inc.
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996

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Device record: <https://www.510kdatabase.net/k772350/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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