

K772178 PACER MONITOR, PULSE WIDTHDec 8, 1977
16 days to decisionK772178 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k772178/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Nov 22, 1977
Decision date	Dec 8, 1977
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Survival Technology, Inc.
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1977-1993

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k772178/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026