

**K873716 ADULT WALKING HEEL PRODUCT NO. 4183-130**Oct 2, 1987  
17 days to decisionK873716 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k873716/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Sep 15, 1987
Decision date	Oct 2, 1987
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Martin Medical</b>
Location	Desoto, KS, US
Contact	RANDY KILBURN
510(k) history	14 submissions · 14 cleared · 1986-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k873716/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 22, 2026