

K895535 PACEVIEWMar 19, 1990
187 days to decisionK895535 · Product code: **KRE** · Cardiovascular
Source: <https://www.510kdatabase.net/k895535/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function, Indirect (KRE)
Date received	Sep 13, 1989
Decision date	Mar 19, 1990
Days to decision	187 days
Third-party review	No

APPLICANT

Company	Micromedical
Location	Gettysburg, PA, US
Contact	WESLEY KULP
510(k) history	2 submissions · 2 cleared · 1990-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895535/>; 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