

**K991007 K-DEFIB/PACE RADIOLUCENT ADULT ELECTRODE,  
MODEL KRT-60**Aug 23, 1999  
150 days to decisionK991007 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k991007/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Mar 26, 1999
Decision date	Aug 23, 1999
Days to decision	150 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Katecho, Inc.</b>
Location	Des Moines, IA, US
Contact	LORNE SCHARNBERG
510(k) history	26 submissions · 25 cleared · 1984-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991007/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 30, 2026