

K800836 KMA FLOW DIRECTED THERMODILUTION CATHETEApr 24, 1980
9 days to decisionK800836 · Product code: **KRB** · CardiovascularSource: <https://www.510kdatabase.net/k800836/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Thermodilution (KRB)
Date received	Apr 15, 1980
Decision date	Apr 24, 1980
Days to decision	9 days
Third-party review	No

APPLICANT

Company	Kma, Inc.
Location	Walker, MI, US
510(k) history	3 submissions · 3 cleared · 1980-1985

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Device record: <https://www.510kdatabase.net/k800836/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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