

K830321 VENTURE II MONITORING SYSTEMJun 3, 1983
123 days to decisionK830321 · Product code: **DSF** · CardiovascularSource: <https://www.510kdatabase.net/k830321/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Jan 31, 1983
Decision date	Jun 3, 1983
Days to decision	123 days
Third-party review	No

APPLICANT

Company	Rae Medical Corp.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1983

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Device record: <https://www.510kdatabase.net/k830321/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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