FDA 510(k) Premarket Notification

K830321 VENTURE II MONITORING SYSTEM

CLEARED

Jun 3, 1983
123 days to decision

K830321 · Product code: **DSF** · Cardiovascular
Source: 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