FDA 510(k) Premarket Notification

K864401 PLURIMUS CARDIAC EVALUATION SYSTEM

CLEARED

Mar 3, 1987
117 days to decision

K864401 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k864401/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Nov 6, 1986
Decision date	Mar 3, 1987
Days to decision	117 days
Third-party review	No

APPLICANT

Company	**Vivix Corp.**
Location	San Jose, CA, US
Contact	L. C BADAGLIACCA
510(k) history	1 submissions · 1 cleared · 1987-1987