FDA 510(k) Premarket Notification

K812914 MICRO-PREPROCESSOR UNIT

CLEARED

Dec 29, 1981
71 days to decision

K812914 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k812914/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 19, 1981
Decision date	Dec 29, 1981
Days to decision	71 days
Third-party review	No

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
510(k) history	254 submissions · 254 cleared · 1976-2011