FDA 510(k) Premarket Notification

K801762 MICRO-PROCESSOR UNIT #A3600AC

CLEARED

Aug 27, 1980
30 days to decision

K801762 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k801762/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 28, 1980
Decision date	Aug 27, 1980
Days to decision	30 days
Third-party review	No

APPLICANT

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