FDA 510(k) Premarket Notification

K770760 MONITOR, PORTABLE, PATIENT, G.E.

CLEARED

May 6, 1977
11 days to decision

K770760 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k770760/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Apr 25, 1977
Decision date	May 6, 1977
Days to decision	11 days
Third-party review	No

APPLICANT

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