

K770760 MONITOR, PORTABLE, PATIENT, G.E.May 6, 1977
11 days to decisionK770760 · Product code: **DPS** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k770760/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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