FDA 510(k) Premarket Notification

K802600 AMP PORTABLE PATIENT PRESSURE MONITOR

CLEARED

Nov 12, 1980
22 days to decision

K802600 · Product code: **DSK** · Cardiovascular
Source: https://www.510kdatabase.net/k802600/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Blood-pressure (DSK)
Date received	Oct 21, 1980
Decision date	Nov 12, 1980
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**American Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1978-1992