

K802600 AMP PORTABLE PATIENT PRESSURE MONITORNov 12, 1980
22 days to decisionK802600 · Product code: **DSK** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k802600/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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