

K800025 DINAMAP RESEARCH MONITOR 1245Feb 1, 1980
25 days to decisionK800025 · Product code: **DXN** · CardiovascularSource: <https://www.510kdatabase.net/k800025/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jan 7, 1980
Decision date	Feb 1, 1980
Days to decision	25 days
Third-party review	No

APPLICANT

Company	Applied Medical Research
Location	Mchenry, IL, US
510(k) history	22 submissions · 22 cleared · 1977-1980

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800025/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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