FDA 510(k) Premarket Notification

K800025 DINAMAP RESEARCH MONITOR 1245

CLEARED

Feb 1, 1980
25 days to decision

K800025 · Product code: **DXN** · Cardiovascular
Source: 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