FDA 510(k) Premarket Notification

K800066 NIASTAT

CLEARED

Feb 29, 1980
49 days to decision

K800066 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k800066/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Jan 11, 1980
Decision date	Feb 29, 1980
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Invivo Research Labs, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1987