

K800066 NIASTATFeb 29, 1980
49 days to decisionK800066 · 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 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800066/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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