FDA 510(k) Premarket Notification

K790123 STIMULATOR, BATTERY POWERED

CLEARED

Feb 6, 1979
14 days to decision

K790123 · Product code: **BXN** · Anesthesiology
Source: https://www.510kdatabase.net/k790123/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Battery-powered (BXN)
Date received	Jan 23, 1979
Decision date	Feb 6, 1979
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Intermedics, Inc.**
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k790123/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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