FDA 510(k) Premarket Notification

K890567 BITE BLOCK #59430

CLEARED

Feb 17, 1989
11 days to decision

K890567 · Product code: **JXL** · Neurology
Source: https://www.510kdatabase.net/k890567/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Block, Bite (JXL)
Date received	Feb 6, 1989
Decision date	Feb 17, 1989
Days to decision	11 days
Third-party review	No

APPLICANT

Company	**Medi-Source, Inc.**
Location	Syosset, NY, US
Contact	ANDREW GALAMBOS
510(k) history	4 submissions · 4 cleared · 1989-1990