

K890567 BITE BLOCK #59430Feb 17, 1989
11 days to decisionK890567 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890567/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026