FDA 510(k) Premarket Notification

K862915 MULTISTIM(4-CHANNEL), MULTISTIM-OCTRODE(8 CHANNEL)

CLEARED

Oct 6, 1986
66 days to decision

K862915 · Product code: **GZB** · Neurology
Source: https://www.510kdatabase.net/k862915/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Aug 1, 1986
Decision date	Oct 6, 1986
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Neuromed, Inc.**
Location	Ft. Lauderdale, FL, US
Contact	FRANK LISKA
510(k) history	15 submissions · 15 cleared · 1985-1996