FDA 510(k) Premarket Notification

K905255 NEUROMED MODEL 1992 LS LAMITRODE LEAD

CLEARED

Jan 25, 1991
65 days to decision

K905255 · Product code: **GZB** · Neurology
Source: https://www.510kdatabase.net/k905255/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Nov 21, 1990
Decision date	Jan 25, 1991
Days to decision	65 days
Third-party review	No

APPLICANT

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