FDA 510(k) Premarket Notification

K896450 NEURO N 50 LESION GENERATOR

CLEARED

Mar 19, 1991
495 days to decision

K896450 · Product code: **GXD** · Neurology
Source: https://www.510kdatabase.net/k896450/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Lesion, Radiofrequency (GXD)
Date received	Nov 9, 1989
Decision date	Mar 19, 1991
Days to decision	495 days
Third-party review	No

APPLICANT

Company	**Leibinger & Fischer , Ltd.**
Location	Dallas, TX, US
Contact	PAUL STEWART
510(k) history	2 submissions · 2 cleared · 1990-1991