FDA 510(k) Premarket Notification

K022222 LAMITRODE S-SERIES (S4 AND S8) LEADS

CLEARED

Aug 8, 2002
30 days to decision

K022222 · Product code: **GZB** · Neurology
Source: https://www.510kdatabase.net/k022222/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Jul 9, 2002
Decision date	Aug 8, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Advanced Neuromodulation System,Inc**
Location	Plano, TX, US
Contact	KATRYNA WARREN
510(k) history	1 submissions · 1 cleared · 2002-2002