FDA 510(k) Premarket Notification

K864956 DYNEX III TRANSCUTANEOUS ELECTRICAL NERVE STIMULA.

CLEARED

Mar 9, 1987
81 days to decision

K864956 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k864956/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 18, 1986
Decision date	Mar 9, 1987
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**La Jolla Technology, Inc.**
Location	Mchenry, IL, US
Contact	BALNAVE, PH.D.
510(k) history	5 submissions · 5 cleared · 1983-1987