FDA 510(k) Premarket Notification

K830932 NYNEX II

CLEARED

Apr 18, 1983
26 days to decision

K830932 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k830932/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 23, 1983
Decision date	Apr 18, 1983
Days to decision	26 days
Third-party review	No

APPLICANT

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