

K830932 NYNEX IIApr 18, 1983
26 days to decisionK830932 · 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

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Device record: <https://www.510kdatabase.net/k830932/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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