FDA 510(k) Premarket Notification

K896620 NEUROTENS

CLEARED

Apr 26, 1990
156 days to decision

K896620 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k896620/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Nov 21, 1989
Decision date	Apr 26, 1990
Days to decision	156 days
Third-party review	No

APPLICANT

Company	**Henley Intl.**
Location	Houston, TX, US
Contact	ERNEST J HENLEY,PHD
510(k) history	44 submissions · 41 cleared · 1986-1995