

K881708 INDICATENS 4X4Aug 11, 1988
113 days to decisionK881708 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k881708/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Apr 20, 1988
Decision date	Aug 11, 1988
Days to decision	113 days
Third-party review	No

APPLICANT

Company	Acudata Software
Location	Petaluma, CA, US
Contact	ROSSEN, DVM
510(k) history	7 submissions · 7 cleared · 1988-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881708/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026