

**K982324 VECTRA PRO MODELS 2 AND 4**Feb 1, 1999  
214 days to decisionK982324 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k982324/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 2, 1998
Decision date	Feb 1, 1999
Days to decision	214 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Chattanooga Group, Inc.</b>
Location	Mchenry, IL, US
Contact	CHERYL G DYER
510(k) history	70 submissions · 68 cleared · 1980-2001

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982324/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026