

**K874735 DANTEC 13L35/36/37 STIM. AND RECORDING ELECTRODE**Jan 22, 1988  
65 days to decisionK874735 · Product code: **GXY** · Neurology  
Source: <https://www.510kdatabase.net/k874735/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Cutaneous (GXY)
Date received	Nov 18, 1987
Decision date	Jan 22, 1988
Days to decision	65 days
Third-party review	No

**APPLICANT**

---

Company	<b>Dantec Electronics, Inc.</b>
Location	Walker, MI, US
Contact	HENRIK HENRIKSEN
510(k) history	12 submissions · 12 cleared · 1985-1990

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874735/> 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