

K880925 DISPOSABLE WET FIELD ERASERApr 13, 1988
40 days to decisionK880925 · Product code: **GXI** · Neurology
Source: <https://www.510kdatabase.net/k880925/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Radiofrequency Lesion (GXI)
Date received	Mar 4, 1988
Decision date	Apr 13, 1988
Days to decision	40 days
Third-party review	No

APPLICANT

Company	Advanced Surgical Products, Inc.
Location	Charles, MO, US
Contact	TODD J HESSEL
510(k) history	64 submissions · 64 cleared · 1987-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880925/>; 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 June 30, 2026