

K871964 SIEGEN NEUROSCOPE SEEG FOR EVOKED RESPONSE STIMU.Oct 1, 1987
134 days to decisionK871964 · Product code: **GWF** · Neurology
Source: <https://www.510kdatabase.net/k871964/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	May 20, 1987
Decision date	Oct 1, 1987
Days to decision	134 days
Third-party review	No

APPLICANT

Company	Siegen Corp.
Location	Mountain View, CA, US
Contact	BERT W MYERS
510(k) history	6 submissions · 6 cleared · 1984-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k871964/>. 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