

K844616 SIEGEN VISUAL STIMULATORFeb 14, 1985
79 days to decisionK844616 · Product code: **GWE** · Neurology
Source: <https://www.510kdatabase.net/k844616/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Nov 27, 1984
Decision date	Feb 14, 1985
Days to decision	79 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k844616/>; 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