FDA 510(k) Premarket Notification

K844616 SIEGEN VISUAL STIMULATOR

CLEARED

Feb 14, 1985
79 days to decision

K844616 · 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