

K781313 VISION-STIMULATOR, CAMJan 26, 1979
179 days to decisionK781313 · Product code: **HPL** · Ophthalmic
Source: <https://www.510kdatabase.net/k781313/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fixation, Ac-powered, Ophthalmic (HPL)
Date received	Jul 31, 1978
Decision date	Jan 26, 1979
Days to decision	179 days
Third-party review	No

APPLICANT

Company	Clement Clarke, Inc.
Location	MD, US
510(k) history	10 submissions · 10 cleared · 1979-1996

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Device record: <https://www.510kdatabase.net/k781313/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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