

K802684 HELIOMATNov 24, 1980
27 days to decisionK802684 · Product code: **EAY** · Dental
Source: <https://www.510kdatabase.net/k802684/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Fiber Optic, Dental (EAY)
Date received	Oct 28, 1980
Decision date	Nov 24, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Vivadent (Usa), Inc.
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1978-1986

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Device record: <https://www.510kdatabase.net/k802684/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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