

K801227 SILICAPAug 12, 1980
81 days to decisionK801227 · Product code: **EMA** · Dental
Source: <https://www.510kdatabase.net/k801227/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Dental (EMA)
Date received	May 23, 1980
Decision date	Aug 12, 1980
Days to decision	81 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/k801227/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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