

K800060 PROVITJan 23, 1980
13 days to decisionK800060 · Product code: **EMA** · Dental
Source: <https://www.510kdatabase.net/k800060/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Dental (EMA)
Date received	Jan 10, 1980
Decision date	Jan 23, 1980
Days to decision	13 days
Third-party review	No

APPLICANT

Company	Parker Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1977-2024

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Device record: <https://www.510kdatabase.net/k800060/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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