FDA 510(k) Premarket Notification

# K801227 SILICAP

**CLEARED**

Aug 12, 1980
81 days to decision

K801227 · 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 |