FDA 510(k) Premarket Notification

# K811582 THE PLACER

**CLEARED**

Jul 28, 1981
53 days to decision

K811582 · Product code: **EID** · Dental
Source: https://www.510kdatabase.net/k811582/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Syringe, Restorative And Impression Material (EID) |
| Date received | Jun 5, 1981 |
| Decision date | Jul 28, 1981 |
| Days to decision | 53 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Cavitron Corp.** |
| Location | Mchenry, IL, US |
| 510(k) history | 32 submissions · 32 cleared · 1976-1981 |