FDA 510(k) Premarket Notification

# K823280 DISTAFLEX

**CLEARED**

Nov 22, 1982
19 days to decision

K823280 · Product code: **EGG** · Dental
Source: https://www.510kdatabase.net/k823280/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Attachment, Precision, All (EGG) |
| Date received | Nov 3, 1982 |
| Decision date | Nov 22, 1982 |
| Days to decision | 19 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Duradent Corp.** |
| Location | Mchenry, IL, US |
| 510(k) history | 1 submissions · 1 cleared · 1982-1982 |