

K823280 DISTAFLEXNov 22, 1982
19 days to decisionK823280 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823280/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026