FDA 510(k) Premarket Notification

K802685 REOLIT

CLEARED

Nov 12, 1980
15 days to decision

K802685 · Product code: **EJK** · Dental
Source: https://www.510kdatabase.net/k802685/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Oct 28, 1980
Decision date	Nov 12, 1980
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Vivadent (Usa), Inc.**
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1978-1986