FDA 510(k) Premarket Notification

K820215 IMPROVED DYCAL

CLEARED

Mar 23, 1982
56 days to decision

K820215 · Product code: **EJK** · Dental
Source: https://www.510kdatabase.net/k820215/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Jan 26, 1982
Decision date	Mar 23, 1982
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Dentsply Intl.**
Location	Walker, MI, US
510(k) history	279 submissions · 279 cleared · 1976-2013