FDA 510(k) Premarket Notification

K781688 LINER, CAVITY CALCIUM HYDROXIDE

CLEARED

Dec 20, 1978
77 days to decision

K781688 · Product code: **EJK** · Dental
Source: https://www.510kdatabase.net/k781688/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Oct 4, 1978
Decision date	Dec 20, 1978
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Kerr Corporation (Danbury)**
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1978-2005