FDA 510(k) Premarket Notification

K872147 KERR CAVALITE

K872147 · Product code: **EJK** · Dental
Source: https://www.510kdatabase.net/k872147/

CLEARED

Aug 26, 1987
83 days to decision

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Jun 4, 1987
Decision date	Aug 26, 1987
Days to decision	83 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k872147/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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