

**K781688 LINER, CAVITY CALCIUM HYDROXIDE**Dec 20, 1978  
77 days to decisionK781688 · Product code: **EJK** · DentalSource: <https://www.510kdatabase.net/k781688/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k781688/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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