

**K802685 REOLIT**Nov 12, 1980  
15 days to decisionK802685 · Product code: **EJK** · DentalSource: <https://www.510kdatabase.net/k802685/>**SUBMISSION DETAILS**

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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**

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

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

Device record: <https://www.510kdatabase.net/k802685/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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