

**K873882 ALKALINER**Feb 26, 1988  
156 days to decisionK873882 · Product code: **EJK** · DentalSource: <https://www.510kdatabase.net/k873882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Liner, Cavity, Calcium Hydroxide (EJK)
Date received	Sep 23, 1987
Decision date	Feb 26, 1988
Days to decision	156 days
Third-party review	No

**APPLICANT**

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Company	<b>Espe GmbH (Us)</b>
Location	Mchenry, IL, US
Contact	HAROLD A OPOTOW
510(k) history	59 submissions · 57 cleared · 1977-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k873882/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 3, 2026