

**K881992 SYRETTE**Jul 26, 1988  
75 days to decisionK881992 · Product code: **EKJ** · DentalSource: <https://www.510kdatabase.net/k881992/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Burnisher, Operative (EKJ)
Date received	May 12, 1988
Decision date	Jul 26, 1988
Days to decision	75 days
Third-party review	No

**APPLICANT**

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Company	<b>Oratec Corp.</b>
Location	Herndon, VA, US
Contact	KELNER, PHD
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881992/>; 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 5, 2026