

**K031681 STERISIL ANTIMICROBIAL TUBING AND BOTTLE**Oct 30, 2003  
153 days to decisionK031681 · Product code: **EIA** · DentalSource: <https://www.510kdatabase.net/k031681/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Operative Dental (EIA)
Date received	May 30, 2003
Decision date	Oct 30, 2003
Days to decision	153 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Sterisil</b>
Location	Castle Rock, CO, US
Contact	BRAD DOWNS
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031681/> 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 1, 2026