

**K013881 RIBBOND - TRIAXIAL**Jan 25, 2002  
63 days to decisionK013881 · Product code: **EBI** · DentalSource: <https://www.510kdatabase.net/k013881/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Nov 23, 2001
Decision date	Jan 25, 2002
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ribbon, Inc.</b>
Location	Seattle, WA, US
Contact	SHOSHANA RUDO DRIVER
510(k) history	2 submissions · 2 cleared · 1991-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013881/> 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 2, 2026