

**K092384 LIQUICORD**Nov 4, 2009  
91 days to decisionK092384 · Product code: **MVL** · Dental  
Source: <https://www.510kdatabase.net/k092384/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cord, Retraction (MVL)
Date received	Aug 5, 2009
Decision date	Nov 4, 2009
Days to decision	91 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Centrix, Inc.</b>
Location	Mchenry, IL, US
Contact	ROGER MASTRONY
510(k) history	47 submissions · 47 cleared · 1979-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092384/> 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 May 14, 2026