FDA 510(k) Premarket Notification

K092384 LIQUICORD

CLEARED

Nov 4, 2009
91 days to decision

K092384 · Product code: **MVL** · Dental
Source: https://www.510kdatabase.net/k092384/

SUBMISSION DETAILS

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

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