FDA 510(k) Premarket Notification

K013881 RIBBOND - TRIAXIAL

CLEARED

Jan 25, 2002
63 days to decision

K013881 · Product code: **EBI** · Dental
Source: https://www.510kdatabase.net/k013881/

SUBMISSION DETAILS

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

Company	**Ribbond, Inc.**
Location	Seattle, WA, US
Contact	SHOSHANA RUDO DRIVER
510(k) history	2 submissions · 2 cleared · 1991-2002