FDA 510(k) Premarket Notification

K190220 Vista FS, Vista FS Liquid

CLEARED

Jun 10, 2019
125 days to decision

K190220 · Product code: **MVL** · Dental
Source: https://www.510kdatabase.net/k190220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cord, Retraction (MVL)
Date received	Feb 5, 2019
Decision date	Jun 10, 2019
Days to decision	125 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Inter-Med/Vista Dental Products**
Location	Racine, WI, US
Contact	Alex Johnson
510(k) history	6 submissions · 6 cleared · 2017-2020