FDA 510(k) Premarket Notification

K210333 DiaPaste

CLEARED

Mar 25, 2021
48 days to decision

K210333 · Product code: **KIF** · Dental
Source: https://www.510kdatabase.net/k210333/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Resin, Root Canal Filling (KIF)
Date received	Feb 5, 2021
Decision date	Mar 25, 2021
Days to decision	48 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**DiaDent Group International**
Location	Chungcheong Buk-Do, KR
Contact	Kab Sun Lee
510(k) history	19 submissions · 19 cleared · 2013-2026