

K210333 DiaPasteMar 25, 2021
48 days to decisionK210333 · Product code: **KIF** · DentalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k210333/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026