

**K182009 Dia-Proseal**Sep 26, 2019  
426 days to decisionK182009 · Product code: **KIF** · Dental  
Source: <https://www.510kdatabase.net/k182009/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Root Canal Filling (KIF)
Date received	Jul 27, 2018
Decision date	Sep 26, 2019
Days to decision	426 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>DiaDent Group International</b>
Location	Chungcheong Buk-Do, KR
Contact	Myung Sub Kim
510(k) history	19 submissions · 19 cleared · 2013-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k182009/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026