

**K150392 PROPEL device**Sep 18, 2015  
212 days to decisionK150392 · Product code: **DZJ** · DentalSource: <https://www.510kdatabase.net/k150392/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Driver, Wire, And Bone Drill, Manual (DZJ)
Date received	Feb 18, 2015
Decision date	Sep 18, 2015
Days to decision	212 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Propel Orthodontics, LLC</b>
Location	Milpitas, CA, US
Contact	BRYCE WAY
510(k) history	3 submissions · 3 cleared · 2015-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k150392/> 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 June 28, 2026