FDA 510(k) Premarket Notification

K150392 PROPEL device

CLEARED

Sep 18, 2015
212 days to decision

K150392 · Product code: **DZJ** · Dental
Source: https://www.510kdatabase.net/k150392/

SUBMISSION DETAILS

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

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