

**K030146 KAVO KEY LASER 1242 AND KAVO KEY LASER  
1243+**Aug 3, 2005  
931 days to decisionK030146 · Product code: **NVK** · Dental  
Source: <https://www.510kdatabase.net/k030146/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Dental, Soft Tissue (NVK)
Date received	Jan 15, 2003
Decision date	Aug 3, 2005
Days to decision	931 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kavo America</b>
Location	Washington, DC, US
Contact	JOHN R FRANZ
510(k) history	20 submissions · 19 cleared · 1991-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030146/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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