

**K130130 ATG/SM-OSA APPLIANCES**Aug 26, 2013  
221 days to decisionK130130 · Product code: **LQZ** · Dental  
Source: <https://www.510kdatabase.net/k130130/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Jaw Repositioning (LQZ)
Date received	Jan 17, 2013
Decision date	Aug 26, 2013
Days to decision	221 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Selane Products, Inc.</b>
Location	Bradenton, FL, US
Contact	CAROLYN PRIMUS
510(k) history	2 submissions · 2 cleared · 2013-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130130/> 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 1, 2026