

**K090911 IST SNORING APPLIANCE**Sep 11, 2009  
163 days to decisionK090911 · Product code: **LQZ** · Dental  
Source: <https://www.510kdatabase.net/k090911/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Jaw Repositioning (LQZ)
Date received	Apr 1, 2009
Decision date	Sep 11, 2009
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dental Crafters</b>
Location	Marshfeld, WI, US
Contact	AMY NYSTROM
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090911/> 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 22, 2026