

**K971818 SNOREX**Dec 18, 1997  
216 days to decisionK971818 · Product code: **LQZ** · DentalSource: <https://www.510kdatabase.net/k971818/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Jaw Repositioning (LQZ)
Date received	May 16, 1997
Decision date	Dec 18, 1997
Days to decision	216 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Snorex (Nz) , Ltd.</b>
Location	Springfield, VA, US
Contact	VERNON PRIBBLE
510(k) history	1 submissions · 1 cleared · 1997-1997

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k971818/> 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 July 2, 2026