FDA 510(k) Premarket Notification

K971818 SNOREX

CLEARED

Dec 18, 1997
216 days to decision

K971818 · Product code: **LQZ** · Dental
Source: 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	**Snorex (Nz) , Ltd.**
Location	Springfield, VA, US
Contact	VERNON PRIBBLE
510(k) history	1 submissions · 1 cleared · 1997-1997