FDA 510(k) Premarket Notification

K011723 ANTI-SNORING DEVICE OR ASD

CLEARED

Dec 18, 2002
562 days to decision

K011723 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k011723/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Jun 4, 2001
Decision date	Dec 18, 2002
Days to decision	562 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pi Medical**
Location	Minneapolis, MN, US
Contact	LINDA S ALEXANDER
510(k) history	1 submissions · 1 cleared · 2002-2002