FDA 510(k) Premarket Notification

K932932 SNOREMASTER SNORE REMEDY

CLEARED

Feb 10, 1995
604 days to decision

K932932 · Product code: **LQZ** · Dental
Source: https://www.510kdatabase.net/k932932/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Jaw Repositioning (LQZ)
Date received	Jun 16, 1993
Decision date	Feb 10, 1995
Days to decision	604 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Snoremaster Co.**
Location	Philadelphia, PA, US
Contact	LEROY S FORNEY
510(k) history	1 submissions · 1 cleared · 1995-1995