FDA 510(k) Premarket Notification

K234089 DNA Appliance

CLEARED

Sep 16, 2024
269 days to decision

K234089 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k234089/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Dec 22, 2023
Decision date	Sep 16, 2024
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vivos Therapeutics**
Location	Murray, KY, US
Contact	Kimberly Griffith
510(k) history	1 submissions · 1 cleared · 2024-2024