FDA 510(k) Premarket Notification

K992694 MODIFICATION TO K6-I DIAGNOSTIC SYSTEM, MODEL K6-I

CLEARED

Sep 10, 1999
29 days to decision

K992694 · Product code: **KZM** · Dental
Source: https://www.510kdatabase.net/k992694/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Muscle Monitoring (KZM)
Date received	Aug 12, 1999
Decision date	Sep 10, 1999
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Myotronics-Noromed, Inc.**
Location	Tukwila, WA, US
Contact	FRAY ADIB
510(k) history	7 submissions · 7 cleared · 1999-2011