

**K992694 MODIFICATION TO K6-I DIAGNOSTIC SYSTEM,
MODEL K6-I**Sep 10, 1999
29 days to decisionK992694 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992694/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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