FDA 510(k) Premarket Notification

K922029 REMUSK MP

CLEARED

Oct 6, 1992
158 days to decision

K922029 · Product code: **LBB** · Neurology
Source: https://www.510kdatabase.net/k922029/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dynamometer, Ac-powered (LBB)
Date received	May 1, 1992
Decision date	Oct 6, 1992
Days to decision	158 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Equipement Biomedical Specialise, Inc.**
Location	Canada, CA
Contact	SYLVAIN BOUCHER
510(k) history	1 submissions · 1 cleared · 1992-1992