FDA 510(k) Premarket Notification

K925225 THE BTE VECTOR

CLEARED

Jun 27, 1994
619 days to decision

K925225 · Product code: **KQX** · Neurology
Source: https://www.510kdatabase.net/k925225/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Oct 16, 1992
Decision date	Jun 27, 1994
Days to decision	619 days
Third-party review	No
Summary / Statement	Summary
Other names	ANALYSIS SYSTEM FOR THE LOW BACK

APPLICANT

Company	**Baltimore Therapeutic Equipment Co.**
Location	Mchenry, IL, US
Contact	KENT FIFE
510(k) history	7 submissions · 7 cleared · 1980-1995