

K925225 THE BTE VECTORJun 27, 1994
619 days to decisionK925225 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925225/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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