FDA 510(k) Premarket Notification

K812908 ELECTRO-GONIOMETER

CLEARED

Nov 24, 1981
36 days to decision

K812908 · Product code: **KQX** · Neurology
Source: https://www.510kdatabase.net/k812908/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Goniometer, Ac-powered (KQX)
Date received	Oct 19, 1981
Decision date	Nov 24, 1981
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Chattanooga Group, Inc.**
Location	Mchenry, IL, US
510(k) history	70 submissions · 68 cleared · 1980-2001