FDA 510(k) Premarket Notification

K782036 TREMOR MONITOR

CLEARED

Jan 23, 1979
49 days to decision

K782036 · Product code: **GYD** · Neurology
Source: https://www.510kdatabase.net/k782036/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Tremor (GYD)
Date received	Dec 5, 1978
Decision date	Jan 23, 1979
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Columbus Instruments Intl. Corp.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1990