FDA 510(k) Premarket Notification

K063872 KINESIA

CLEARED

Apr 6, 2007
98 days to decision

K063872 · Product code: **GYD** · Neurology
Source: https://www.510kdatabase.net/k063872/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Tremor (GYD)
Date received	Dec 29, 2006
Decision date	Apr 6, 2007
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cleveland Medical Devices, Inc.**
Location	Cleveland, OH, US
Contact	JOSEPH GIUFFRIDA
510(k) history	8 submissions · 8 cleared · 1995-2007