FDA 510(k) Premarket Notification

K843923 THYMATRON

CLEARED

Dec 3, 1984
59 days to decision

K843923 · Product code: **QGH** · Neurology
Source: https://www.510kdatabase.net/k843923/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electroconvulsive Therapy Device For Catatonia, Major Depressive Disorder, And Bipolar Disorder (QGH)
Date received	Oct 5, 1984
Decision date	Dec 3, 1984
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Somatics, Inc.**
Location	Lake Bluff, IL, US
Contact	RICHARD ABRAMS
510(k) history	7 submissions · 7 cleared · 1984-1999