FDA 510(k) Premarket Notification

K860467 ELECTROSHOCK UNIT NEUROLOGY MODEL B-25

CLEARED

Nov 10, 1986
277 days to decision

K860467 · Product code: **QGH** · Neurology
Source: https://www.510kdatabase.net/k860467/

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	Feb 6, 1986
Decision date	Nov 10, 1986
Days to decision	277 days
Third-party review	No

APPLICANT

Company	**Medcraft Corp.**
Location	Mchenry, IL, US
Contact	CHYUNG
510(k) history	5 submissions · 5 cleared · 1976-1986