FDA 510(k) Premarket Notification

K960754 SPECTRUM 5000 Q, 5000 M, 4000 Q, 4000 M

CLEARED

Sep 18, 1996
208 days to decision

K960754 · Product code: **QGH** · Neurology
Source: https://www.510kdatabase.net/k960754/

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 23, 1996
Decision date	Sep 18, 1996
Days to decision	208 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mecta Corp.**
Location	Portland, OR, US
Contact	ROBIN H NICOL
510(k) history	3 submissions · 3 cleared · 1985-1997