FDA 510(k) Premarket Notification

# K911144 MF-500, MODIFICATION

**CLEARED**

Oct 18, 1991
235 days to decision

K911144 · Product code: **QGH** · Neurology
Source: https://www.510kdatabase.net/k911144/

## 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 25, 1991 |
| Decision date | Oct 18, 1991 |
| Days to decision | 235 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Elcot, Inc.** |
| Location | New York, NY, US |
| Contact | IVAN G SCHICK |
| 510(k) history | 2 submissions · 2 cleared · 1987-1991 |