

**K911144 MF-500, MODIFICATION**Oct 18, 1991  
235 days to decisionK911144 · Product code: **QGH** · Neurology  
Source: <https://www.510kdatabase.net/k911144/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911144/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 1, 2026