

**K843923 THYMATRON**Dec 3, 1984  
59 days to decisionK843923 · Product code: **QGH** · Neurology  
Source: <https://www.510kdatabase.net/k843923/>**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	Oct 5, 1984
Decision date	Dec 3, 1984
Days to decision	59 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843923/>; 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 June 30, 2026