

**K820414 UNIVERSAL TESTING & ANALYSIS SYSTEMS**Mar 4, 1982  
20 days to decisionK820414 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k820414/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Feb 12, 1982
Decision date	Mar 4, 1982
Days to decision	20 days
Third-party review	No

**APPLICANT**

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Company	<b>Lkc Technologies, Inc.</b>
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k820414/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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