

**K960591 RE-USABLE BIPOLAR CONCENTRIC  
NEEDLE(237-XXX-24,237-XXX-24STP,  
237-XXX-24TP,237-XXX-24STP**Jun 13, 1996  
122 days to decisionK960591 · Product code: **IKT** · Neurology  
Source: <https://www.510kdatabase.net/k960591/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Needle, Diagnostic Electromyograph (IKT)
Date received	Feb 12, 1996
Decision date	Jun 13, 1996
Days to decision	122 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Chalgren Enterprises, Inc.</b>
Location	Gilroy, CA, US
Contact	RICHARD KAISER
510(k) history	21 submissions · 21 cleared · 1989-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960591/> 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 29, 2026