

**K834477 DISPOS. SCOTT CANNULAS 16-1054 ETC.**Mar 19, 1984  
90 days to decisionK834477 · Product code: **HCA** · Neurology  
Source: <https://www.510kdatabase.net/k834477/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Ventricular (HCA)
Date received	Dec 20, 1983
Decision date	Mar 19, 1984
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
510(k) history	152 submissions · 151 cleared · 1976-2020

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

Device record: <https://www.510kdatabase.net/k834477/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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