

**K823790 VENTRICULAR ACCESS SYSTEM**Apr 28, 1983  
133 days to decisionK823790 · Product code: **LKG** · General Hospital  
Source: <https://www.510kdatabase.net/k823790/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intraventricular (LKG)
Date received	Dec 16, 1982
Decision date	Apr 28, 1983
Days to decision	133 days
Third-party review	No

**APPLICANT**

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Company	<b>American Heyer Schulte</b>
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823790/> 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 May 21, 2026