

**K833672 HH SUBCUTANEOUS ACCESS PORT**Mar 6, 1984  
140 days to decisionK833672 · Product code: **LKG** · General Hospital  
Source: <https://www.510kdatabase.net/k833672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intraventricular (LKG)
Date received	Oct 18, 1983
Decision date	Mar 6, 1984
Days to decision	140 days
Third-party review	No

**APPLICANT**

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Company	<b>Holter-Hausner Intl.</b>
Location	Mchenry, IL, US
510(k) history	42 submissions · 38 cleared · 1978-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833672/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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