

**K830970 DOUBLE LUMEN TRANSFER DEVICE**Apr 12, 1983  
15 days to decisionK830970 · Product code: **LHI** · General HospitalSource: <https://www.510kdatabase.net/k830970/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Mar 28, 1983
Decision date	Apr 12, 1983
Days to decision	15 days
Third-party review	No

**APPLICANT**

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Company	<b>Codan Trading Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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

Device record: <https://www.510kdatabase.net/k830970/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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