

**K832312 I.V. TRANSFER NEEDLE**Sep 12, 1983  
60 days to decision

K832312 · Product code: LHI · General Hospital

Source: <https://www.510kdatabase.net/k832312/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Jul 14, 1983
Decision date	Sep 12, 1983
Days to decision	60 days
Third-party review	No

**APPLICANT**

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Company	<b>Baxa Corp., Sub. of Cook Group, Inc.</b>
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1978-1991

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

Device record: <https://www.510kdatabase.net/k832312/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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