

**K833705 BETAJECT BD U100**May 9, 1984  
204 days to decisionK833705 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k833705/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 18, 1983
Decision date	May 9, 1984
Days to decision	204 days
Third-party review	No

**APPLICANT**

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Company	<b>Orange Medical Instruments, Inc.</b>
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1982-1986

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

Device record: <https://www.510kdatabase.net/k833705/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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