

**K930217 BAXTER DISPOSABLE INTRAOSSEOUS INFUSION  
NEEDLE**Feb 28, 1994  
409 days to decisionK930217 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k930217/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 15, 1993
Decision date	Feb 28, 1994
Days to decision	409 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Baxter Healthcare Corp</b>
Location	Mchenry, IL, US
Contact	JAN ROBERTSON
510(k) history	505 submissions · 496 cleared · 1977-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930217/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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