

**K991904 MODIFICATION OF JMS NEEDLES AND JMS SYRINGES**Aug 31, 1999  
88 days to decisionK991904 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k991904/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 4, 1999
Decision date	Aug 31, 1999
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Jms Co., Ltd.</b>
Location	Hiroshima, JP
Contact	KEISUKE URATOMI
510(k) history	24 submissions · 24 cleared · 1985-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991904/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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