

**K011925 HYPODERMIC NEEDLE-PRO NEEDLE WITH NEEDLE PROTECTION DEVICE**

Jul 12, 2001  
22 days to decision

K011925 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k011925/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 20, 2001
Decision date	Jul 12, 2001
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary
Other names	HYPODERMIC NEEDLE-PRO SYRINGE & NEEDLE WITH NEEDLE PROTECTIO

**APPLICANT**

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Company	<b>Sims Portex, Inc.</b>
Location	Keene, NH, US
Contact	BRAIN D FARIAS
510(k) history	12 submissions · 12 cleared · 1998-2001

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

Device record: <https://www.510kdatabase.net/k011925/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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