

**K945273 PRO-TECH DISPOSABLE SHARPS CONTAINERS  
VNC2**Dec 14, 1994  
47 days to decisionK945273 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k945273/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 28, 1994
Decision date	Dec 14, 1994
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pro-Tec Containers, Inc.</b>
Location	Irvine, CA, US
Contact	TREESA SPENCER
510(k) history	6 submissions · 6 cleared · 1989-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k945273/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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