

**K945275 PRO-TEC DISPOSBLE SHARPS CONTAINERS Q1**Dec 14, 1994  
47 days to decisionK945275 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k945275/>**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/k945275/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026