

**K894372 NIPRO NEEDLE HOLDER**Sep 28, 1989  
73 days to decisionK894372 · Product code: **FMI** · ChemistrySource: <https://www.510kdatabase.net/k894372/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 17, 1989
Decision date	Sep 28, 1989
Days to decision	73 days
Third-party review	No

**APPLICANT**

---

Company	<b>Pharma-Plast USA, Inc.</b>
Location	Canada H3a 2n4, CA
Contact	LINDEMANN, BOC
510(k) history	9 submissions · 9 cleared · 1985-1989

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894372/>, 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 May 27, 2026