FDA 510(k) Premarket Notification

K894372 NIPRO NEEDLE HOLDER

CLEARED

Sep 28, 1989
73 days to decision

K894372 · Product code: **FMI** · Chemistry
Source: 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	**Pharma-Plast USA, Inc.**
Location	Canada H3a 2n4, CA
Contact	LINDEMANN, BOC
510(k) history	9 submissions · 9 cleared · 1985-1989