

**K903490 NORPLANT(R) TROCAR**Nov 28, 1990  
127 days to decisionK903490 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k903490/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jul 24, 1990
Decision date	Nov 28, 1990
Days to decision	127 days
Third-party review	No

**APPLICANT**

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Company	<b>Wyeth-Ayerst Laboratories, Inc.</b>
Location	Philadelphia, PA, US
Contact	JUSTIN R VICTORIA
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903490/>; 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 July 7, 2026