FDA 510(k) Premarket Notification

K903490 NORPLANT(R) TROCAR

CLEARED

Nov 28, 1990
127 days to decision

K903490 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k903490/

SUBMISSION DETAILS

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

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