

K942159 NOVOPEN 1.5Apr 26, 1995
358 days to decisionK942159 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k942159/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	May 3, 1994
Decision date	Apr 26, 1995
Days to decision	358 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Novo Nordisk Pharmaceuticals, Inc.
Location	Princeton, NJ, US
Contact	GEORGE A CLAY, PH.D
510(k) history	2 submissions · 2 cleared · 1993-1995

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