FDA 510(k) Premarket Notification

K982842 HUMAPEN AND HUMAPEN ERGO

CLEARED

Sep 25, 1998
44 days to decision

K982842 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k982842/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 12, 1998
Decision date	Sep 25, 1998
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Eli Lilly and Co.**
Location	Indianapolis, IN, US
Contact	LEEANN CHAMBERS
510(k) history	2 submissions · 2 cleared · 1998-2007