

K982842 HUMAPEN AND HUMAPEN ERGOSep 25, 1998
44 days to decisionK982842 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982842/> Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026