

K142518 HumaPen LuxuraJun 5, 2015
270 days to decisionK142518 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k142518/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Sep 8, 2014
Decision date	Jun 5, 2015
Days to decision	270 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Eli Lilly and Company
Location	Indianapolis, IN, US
Contact	LeeAnn Chambers, MS., RAC
Website	http://www.lilly.com
510(k) history	5 submissions · 5 cleared · 2006-2022

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