FDA 510(k) Premarket Notification

K113673 MYSAFETY SYRINGE

CLEARED

Aug 3, 2012
234 days to decision

K113673 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k113673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Dec 13, 2011
Decision date	Aug 3, 2012
Days to decision	234 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medicalchain Internartional Corporation**
Location	Thousand Oaks, CA, US
Contact	ROBIN HWANG
510(k) history	1 submissions · 1 cleared · 2012-2012