FDA 510(k) Premarket Notification

CLEARED

K122416 ARTIGLASS L. O.R. GLASS SYRINGE

May 3, 2013
268 days to decision

K122416 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k122416/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 8, 2012
Decision date	May 3, 2013
Days to decision	268 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Artiglass Srl**
Location	Due Carrare Padova, IT
Contact	FEDERICO BACCARIN
510(k) history	2 submissions · 2 cleared · 2013-2022