FDA 510(k) Premarket Notification

K043261 HARVEST GRAFT DELIVERY SYRINGE

CLEARED

Mar 11, 2005
107 days to decision

K043261 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k043261/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Nov 24, 2004
Decision date	Mar 11, 2005
Days to decision	107 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Harvest Technologies, Corp.**
Location	Plymouth, MA, US
Contact	JACK BONASERA
510(k) history	6 submissions · 6 cleared · 2001-2012