FDA 510(k) Premarket Notification

K020252 SMARTJET GRAFTING LIQUID APPLICATOR, MODELS SK/S & LK/2

CLEARED

Apr 5, 2002
71 days to decision

K020252 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k020252/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 24, 2002
Decision date	Apr 5, 2002
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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