FDA 510(k) Premarket Notification

K020666 DUPLOJECT EASY-PREP SYSTEM 1ML, 2ML, 5ML

CLEARED

Jun 24, 2002
115 days to decision

K020666 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k020666/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Syringe, Piston (FMF)
Date received	Mar 1, 2002
Decision date	Jun 24, 2002
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Baxter Healthcare Corp**
Location	Mchenry, IL, US
Contact	ARLENE VIDOR
510(k) history	505 submissions · 496 cleared · 1977-2019