FDA 510(k) Premarket Notification

K031394 AMERICAN SYRINGE COMPANY 1CC AND 3CC SYRINGE

CLEARED

Jul 25, 2003
84 days to decision

K031394 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k031394/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	May 2, 2003
Decision date	Jul 25, 2003
Days to decision	84 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**American Syringe Company**
Location	Miami, FL, US
Contact	DAVID M GARVIN
510(k) history	1 submissions · 1 cleared · 2003-2003