FDA 510(k) Premarket Notification

K960049 BRANNON PORTSYRINGE

CLEARED

May 24, 1996
143 days to decision

K960049 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k960049/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 2, 1996
Decision date	May 24, 1996
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**James K. Brannon**
Location	Culver City, CA, US
Contact	JAMES K BRANNON
510(k) history	2 submissions · 2 cleared · 1996-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k960049/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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