FDA 510(k) Premarket Notification

K941453 CHECK VALVE SOLUTION SET

CLEARED

Aug 16, 1994
144 days to decision

K941453 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k941453/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 25, 1994
Decision date	Aug 16, 1994
Days to decision	144 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Douglas Medical Products Corp.**
Location	Mundelein, IL, US
Contact	DOUGLAS JOHNSON
510(k) history	19 submissions · 19 cleared · 1992-1996