FDA 510(k) Premarket Notification

K921391 SOLUTION ADMINISTRATION SET

CLEARED

Sep 2, 1992
163 days to decision

K921391 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k921391/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 23, 1992
Decision date	Sep 2, 1992
Days to decision	163 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