FDA 510(k) Premarket Notification

K915390 3 LEAD EXTENSION WITH 2 INTERLINK INJECTION SITES

CLEARED

Mar 12, 1992
104 days to decision

K915390 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k915390/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 29, 1991
Decision date	Mar 12, 1992
Days to decision	104 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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