FDA 510(k) Premarket Notification

K911868 INTERLINK(TM) BLUNT CANNULA, MODIFICATION

CLEARED

Jun 24, 1991
59 days to decision

K911868 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k911868/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 26, 1991
Decision date	Jun 24, 1991
Days to decision	59 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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