

K911868 INTERLINK(TM) BLUNT CANNULA, MODIFICATIONJun 24, 1991
59 days to decisionK911868 · 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

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