

**K880482 SECONDARY SOLUTION ADMINISTRATION SET
W/SHEATH**Mar 18, 1988
42 days to decisionK880482 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k880482/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 5, 1988
Decision date	Mar 18, 1988
Days to decision	42 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880482/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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