

K961225 CONTINUFLO SOLUTION SET/SECONDARY MEDICATION SET

Jun 21, 1996
84 days to decision

K961225 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k961225/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 29, 1996
Decision date	Jun 21, 1996
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k961225/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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