

K123868 INTERLINK SYSTEM LEVER LOCK CANNULA WITH CHECK VALVE, SECONDARY MEDICATION SETS, SOLUTION SETS, CONTINU-FLO SOLUTION SET

Jan 8, 2013
22 days to decision

K123868 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k123868/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 17, 2012
Decision date	Jan 8, 2013
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k123868/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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