

K883638 SAFESITE INJECTION SITE AND BLUNT CANNULASep 23, 1988
29 days to decisionK883638 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k883638/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 25, 1988
Decision date	Sep 23, 1988
Days to decision	29 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/k883638/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026