

K933852 PRO-LOK EXTENSION SETDec 29, 1993
145 days to decisionK933852 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k933852/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 6, 1993
Decision date	Dec 29, 1993
Days to decision	145 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Beech Medical Products, Inc.
Location	Newtown, PA, US
Contact	JOHN ROMANO
510(k) history	4 submissions · 4 cleared · 1993-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933852/> 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 July 6, 2026