FDA 510(k) Premarket Notification

K933852 PRO-LOK EXTENSION SET

CLEARED

Dec 29, 1993
145 days to decision

K933852 · 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