FDA 510(k) Premarket Notification

K020419 V SET

CLEARED

Aug 29, 2003
568 days to decision

K020419 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k020419/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 7, 2002
Decision date	Aug 29, 2003
Days to decision	568 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**P.T. Greenleaf**
Location	Perth, AU
Contact	GEORGE O'NEIL
510(k) history	2 submissions · 2 cleared · 2003-2003