FDA 510(k) Premarket Notification

K930926 DURHAM

CLEARED

Oct 12, 1993
231 days to decision

K930926 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k930926/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 23, 1993
Decision date	Oct 12, 1993
Days to decision	231 days
Third-party review	No
Summary / Statement	Statement
Other names	Y CHECK SET-LUER LOCK W/INJECTION SITE

APPLICANT

Company	**Intl. Medical Consultants**
Location	Washington Crossing, PA, US
Contact	JOHN ROMANO
510(k) history	6 submissions · 6 cleared · 1993-1993