FDA 510(k) Premarket Notification

K834071 LYMPHANGIOGRAPHY SET SPECIAL 30G

CLEARED

Jan 25, 1984
61 days to decision

K834071 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k834071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 25, 1983
Decision date	Jan 25, 1984
Days to decision	61 days
Third-party review	No

APPLICANT

Company	**Hyperion, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1984-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k834071/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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