FDA 510(k) Premarket Notification

K781637 MODEL IV 1500 STANDARD BURETTE

CLEARED

Dec 12, 1978
82 days to decision

K781637 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k781637/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 21, 1978
Decision date	Dec 12, 1978
Days to decision	82 days
Third-party review	No

APPLICANT

Company	**Valleylab, Inc.**
Location	Mchenry, IL, US
510(k) history	94 submissions · 93 cleared · 1976-2003