FDA 510(k) Premarket Notification

K800600 B-D STASET, I.V. FLOW CONTROLLER

CLEARED

Apr 16, 1980
30 days to decision

K800600 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k800600/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 17, 1980
Decision date	Apr 16, 1980
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Bd Becton Dickinson Vacutainer Systems Preanalytic**
Location	Washington, DC, US
510(k) history	632 submissions · 625 cleared · 1976-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k800600/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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