FDA 510(k) Premarket Notification

K823406 GUARDIAN 110, VOL. CONTROL DELIV. SYS

CLEARED

Dec 22, 1982
37 days to decision

K823406 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k823406/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Nov 15, 1982
Decision date	Dec 22, 1982
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Avi, Inc.**
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1981-1988