

K823406 GUARDIAN 110, VOL. CONTROL DELIV. SYSDec 22, 1982
37 days to decisionK823406 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823406/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 9, 2026