FDA 510(k) Premarket Notification

K811933 GUARDIAN, VOLUMETRIC CONTROL DELIVERY

CLEARED

Jul 20, 1981
14 days to decision

K811933 · Product code: **FRN** · General Hospital
Source: https://www.510kdatabase.net/k811933/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Jul 6, 1981
Decision date	Jul 20, 1981
Days to decision	14 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/k811933/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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