FDA 510(k) Premarket Notification

K830448 SECURA SET VOLUME LIMITING SET

CLEARED

Feb 25, 1983
10 days to decision

K830448 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k830448/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 15, 1983
Decision date	Feb 25, 1983
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Diatek, Inc.**
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1979-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k830448/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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