FDA 510(k) Premarket Notification

K830377 SECURA SET PRIMARY ADMIN. SET

CLEARED

Mar 1, 1983
25 days to decision

K830377 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k830377/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 4, 1983
Decision date	Mar 1, 1983
Days to decision	25 days
Third-party review	No

APPLICANT

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