FDA 510(k) Premarket Notification

K890109 SAFEPETTE DISPENSER

CLEARED

Feb 23, 1989
44 days to decision

K890109 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k890109/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Jan 10, 1989
Decision date	Feb 23, 1989
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**Safe-Tec Clinical Products, Inc.**
Location	Ivyland, PA, US
Contact	G KENDRICK
510(k) history	2 submissions · 2 cleared · 1988-1989