FDA 510(k) Premarket Notification

K980839 AVOCET PT

CLEARED

Sep 30, 1998
210 days to decision

K980839 · Product code: **JPA** · Hematology
Source: https://www.510kdatabase.net/k980839/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Mar 4, 1998
Decision date	Sep 30, 1998
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Avocet Medical, Inc.**
Location	Campbell, CA, US
Contact	Judith Blunt
510(k) history	4 submissions · 4 cleared · 1998-2000