FDA 510(k) Premarket Notification

K870687 COAGULATION CONTROL, LEVEL III

CLEARED

Apr 20, 1987
60 days to decision

K870687 · Product code: **JPA** · Hematology
Source: https://www.510kdatabase.net/k870687/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Feb 19, 1987
Decision date	Apr 20, 1987
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**U. S. Diagnostics, Inc.**
Location	San Luis Obispo, CA, US
Contact	ROY E SPECK
510(k) history	22 submissions · 22 cleared · 1986-2011