FDA 510(k) Premarket Notification

K895844 THROMBELASTOGRAPH(R)

CLEARED

Oct 24, 1989
21 days to decision

K895844 · Product code: **GKP** · Hematology
Source: https://www.510kdatabase.net/k895844/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Coagulation, Automated (GKP)
Date received	Oct 3, 1989
Decision date	Oct 24, 1989
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Haemoscope Corp.**
Location	Morton Grove, IL, US
Contact	ZUCKERMAN, PHD
510(k) history	7 submissions · 7 cleared · 1989-2007