FDA 510(k) Premarket Notification

K852492 COAGULATOR

CLEARED

Aug 13, 1985
62 days to decision

K852492 · Product code: **KQG** · Hematology
Source: https://www.510kdatabase.net/k852492/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Coagulation (KQG)
Date received	Jun 12, 1985
Decision date	Aug 13, 1985
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Med USA**
Location	Waltham, MA, US
Contact	ISMAIL NABIH
510(k) history	5 submissions · 5 cleared · 1985-1990

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k852492/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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