FDA 510(k) Premarket Notification

K882481 HEMOCHRON CONTROL ACT TEST TUBE NORMAL/ABNOR. I/II

CLEARED

Aug 30, 1988
75 days to decision

K882481 · Product code: **GGN** · Hematology
Source: https://www.510kdatabase.net/k882481/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plasma, Coagulation Control (GGN)
Date received	Jun 16, 1988
Decision date	Aug 30, 1988
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**International Technidyne Corp.**
Location	Walker, MI, US
Contact	LES HEIMANN
510(k) history	47 submissions · 47 cleared · 1983-2013