FDA 510(k) Premarket Notification

K842286 DIA T1, T2 & T3

CLEARED

Sep 11, 1984
90 days to decision

K842286 · Product code: **JPK** · Hematology
Source: https://www.510kdatabase.net/k842286/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	Jun 13, 1984
Decision date	Sep 11, 1984
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Diatech, Inc.**
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1984-1988