FDA 510(k) Premarket Notification

K810719 REPLICA PLATER

CLEARED

Apr 1, 1981
15 days to decision

K810719 · Product code: **JTC** · Hematology
Source: https://www.510kdatabase.net/k810719/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Microtiter Diluting/dispensing (JTC)
Date received	Mar 17, 1981
Decision date	Apr 1, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Dynatech Corp.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1979-1991