FDA 510(k) Premarket Notification

K780305 CELL

CLEARED

Mar 27, 1978
32 days to decision

K780305 · Product code: **KJF** · Pathology
Source: https://www.510kdatabase.net/k780305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suspension, Cell Culture (KJF)
Date received	Feb 23, 1978
Decision date	Mar 27, 1978
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Dynatech Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	44 submissions · 44 cleared · 1978-1996