

K780305 CELLMar 27, 1978
32 days to decisionK780305 · 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

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Device record: <https://www.510kdatabase.net/k780305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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