

**K810719 REPLICA PLATER**Apr 1, 1981  
15 days to decisionK810719 · Product code: **JTC** · Hematology  
Source: <https://www.510kdatabase.net/k810719/>**SUBMISSION DETAILS**

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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**

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

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Device record: <https://www.510kdatabase.net/k810719/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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