

K823773 PROVIAL II #006-010-0201Jan 21, 1983
38 days to decisionK823773 · Product code: **KDT** · Pathology
Source: <https://www.510kdatabase.net/k823773/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Specimen Mailer And Storage, Sterile (KDT)
Date received	Dec 14, 1982
Decision date	Jan 21, 1983
Days to decision	38 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k823773/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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