FDA 510(k) Premarket Notification

K823773 PROVIAL II #006-010-0201

CLEARED

Jan 21, 1983
38 days to decision

K823773 · 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