FDA 510(k) Premarket Notification

K770654 UN, UV DELTA TEST ASSAY

CLEARED

Jun 24, 1977
79 days to decision

K770654 · Product code: **CDQ** · Chemistry
Source: https://www.510kdatabase.net/k770654/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Urease And Glutamic Dehydrogenase, Urea Nitrogen (CDQ)
Date received	Apr 6, 1977
Decision date	Jun 24, 1977
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Bd Becton Dickinson Vacutainer Systems Preanalytic**
Location	Washington, DC, US
510(k) history	632 submissions · 625 cleared · 1976-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k770654/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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