FDA 510(k) Premarket Notification

K770400 REAGENT, AMYLASE LIPASE

CLEARED

Apr 26, 1977
55 days to decision

K770400 · Product code: **KHM** · Chemistry
Source: https://www.510kdatabase.net/k770400/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nephelometric, Amylase (KHM)
Date received	Mar 2, 1977
Decision date	Apr 26, 1977
Days to decision	55 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/k770400/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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