FDA 510(k) Premarket Notification

K830363 BACTEC 170 TRYPTIC SOY BROTH W/RESINS

CLEARED

Mar 24, 1983
48 days to decision

K830363 · Product code: **LJF** · Microbiology
Source: https://www.510kdatabase.net/k830363/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Antimicrobial Drug Removal (LJF)
Date received	Feb 4, 1983
Decision date	Mar 24, 1983
Days to decision	48 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/k830363/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026