FDA 510(k) Premarket Notification

K832229 BBL ANAGEL POWDER

CLEARED

Aug 12, 1983
32 days to decision

K832229 · Product code: **KZJ** · Microbiology
Source: https://www.510kdatabase.net/k832229/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Gas Generating (KZJ)
Date received	Jul 11, 1983
Decision date	Aug 12, 1983
Days to decision	32 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/k832229/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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