FDA 510(k) Premarket Notification

K780303 AUTO ISO-FILTER COLLECTION TUBE CAP

CLEARED

Mar 30, 1978
35 days to decision

K780303 · Product code: **GJE** · Chemistry
Source: https://www.510kdatabase.net/k780303/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tray, Blood Collection (GJE)
Date received	Feb 23, 1978
Decision date	Mar 30, 1978
Days to decision	35 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/k780303/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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