FDA 510(k) Premarket Notification

K760041 FILTER, RESPIRATORY (U-MID)

CLEARED

Jul 16, 1976
31 days to decision

K760041 · Product code: **CAH** · General Hospital
Source: https://www.510kdatabase.net/k760041/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Bacterial, Breathing-circuit (CAH)
Date received	Jun 15, 1976
Decision date	Jul 16, 1976
Days to decision	31 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/k760041/ 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