

K760585 U-MID PREFILLED NEBULIZEROct 15, 1976
42 days to decisionK760585 · Product code: **CAF** · AnesthesiologySource: <https://www.510kdatabase.net/k760585/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Sep 3, 1976
Decision date	Oct 15, 1976
Days to decision	42 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/k760585/> 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