FDA 510(k) Premarket Notification

K760585 U-MID PREFILLED NEBULIZER

CLEARED

Oct 15, 1976
42 days to decision

K760585 · Product code: **CAF** · Anesthesiology
Source: 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