FDA 510(k) Premarket Notification

K880411 DEY-VIAL SODIUM CHLORIDE SOLUTION, USP, STERILE

CLEARED

Feb 29, 1988
28 days to decision

K880411 · Product code: **CAF** · Anesthesiology
Source: https://www.510kdatabase.net/k880411/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Feb 1, 1988
Decision date	Feb 29, 1988
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Dey Laboratories, Inc.**
Location	Mchenry, IL, US
Contact	RAFF, PHD
510(k) history	6 submissions · 6 cleared · 1983-1997