

**K880411 DEY-VIAL SODIUM CHLORIDE SOLUTION, USP,
STERILE**Feb 29, 1988
28 days to decisionK880411 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880411/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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