FDA 510(k) Premarket Notification

CLEARED

K883593 #400 BROOK AIRWAY & #900 PROFESSIONAL BROOK AIRWAY

Sep 30, 1988
38 days to decision

K883593 · Product code: **CAE** · Anesthesiology
Source: https://www.510kdatabase.net/k883593/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Airway, Oropharyngeal, Anesthesiology (CAE)
Date received	Aug 23, 1988
Decision date	Sep 30, 1988
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Ecolab, Inc.**
Location	St. Paul, MN, US
Contact	ANN M OXFORD
Website	http://www.ecolab.com/
510(k) history	7 submissions · 7 cleared · 1988-2019

Ecolab, Inc. is a global provider of water, hygiene, and infection prevention solutions and services. The company operates with a manufacturing facility in St. Paul, Minnesota, serving diverse industries including healthcare, food and beverage processing, and hospitality. Ecolab received FDA 510(k) clearances from total submissions between 1988 and 2019. The company's cleared devices span multiple healthcare specialties, including general hospital equipment, gastroenterology and urology devices, and anesthesiology products. The company is currently inactive, with no submi...