FDA 510(k) Premarket Notification

K780359 OXYGEN REGULATOR/FLOWMETER

CLEARED

Apr 4, 1978
29 days to decision

K780359 · Product code: **CAN** · Anesthesiology
Source: https://www.510kdatabase.net/k780359/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Pressure, Gas Cylinder (CAN)
Date received	Mar 6, 1978
Decision date	Apr 4, 1978
Days to decision	29 days
Third-party review	No

APPLICANT

Company	**Erie Mfg., Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978