FDA 510(k) Premarket Notification

K813436 OXYCON

CLEARED

Jan 7, 1982
30 days to decision

K813436 · Product code: **CAW** · Anesthesiology
Source: https://www.510kdatabase.net/k813436/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Dec 8, 1981
Decision date	Jan 7, 1982
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Home Medical Mfg., Co., Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982