FDA 510(k) Premarket Notification

K790520 AIRCARE OXYGEN GENERATOR

CLEARED

Apr 18, 1979
33 days to decision

K790520 · Product code: **CAW** · Anesthesiology
Source: https://www.510kdatabase.net/k790520/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Mar 16, 1979
Decision date	Apr 18, 1979
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Medical Services of America, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k790520/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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