FDA 510(k) Premarket Notification

K904002 GENOX MODEL CT-1 OXYGEN/NITROGEN GENER-DISTR SYST

CLEARED

Sep 5, 1990
7 days to decision

K904002 · Product code: **CAW** · Anesthesiology
Source: https://www.510kdatabase.net/k904002/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Aug 29, 1990
Decision date	Sep 5, 1990
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**Pacific Consolidated Industries**
Location	Orange, CA, US
Contact	LEE W SMITH
510(k) history	2 submissions · 2 cleared · 1990-1991

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k904002/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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