FDA 510(k) Premarket Notification

K895149 NEOCATH 1000 UMBILICAL ARTERY OXYGEN MONITORING SY

CLEARED

Nov 28, 1989
97 days to decision

K895149 · Product code: **CCE** · Anesthesiology
Source: https://www.510kdatabase.net/k895149/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Partial Pressure, Blood-phase, Indwelling (CCE)
Date received	Aug 23, 1989
Decision date	Nov 28, 1989
Days to decision	97 days
Third-party review	No

APPLICANT

Company	**Shiley, Inc.**
Location	Mchenry, IL, US
Contact	ALLAN ABATI
510(k) history	174 submissions · 174 cleared · 1976-1993

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k895149/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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