FDA 510(k) Premarket Notification

K801607 MOM 2

CLEARED

Aug 7, 1980
23 days to decision

K801607 · Product code: **CCL** · Anesthesiology
Source: https://www.510kdatabase.net/k801607/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Gaseous-phase (CCL)
Date received	Jul 15, 1980
Decision date	Aug 7, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Neotronics, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1991