

K801607 MOM 2Aug 7, 1980
23 days to decisionK801607 · Product code: **CCL** · Anesthesiology
Source: <https://www.510kdatabase.net/k801607/>**SUBMISSION DETAILS**

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|-----------------------|--|
| 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 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801607/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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