

K780359 OXYGEN REGULATOR/FLOWMETERApr 4, 1978
29 days to decisionK780359 · Product code: **CAN** · AnesthesiologySource: <https://www.510kdatabase.net/k780359/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Regulator, Pressure, Gas Cylinder (CAN) |
| Date received | Mar 6, 1978 |
| Decision date | Apr 4, 1978 |
| Days to decision | 29 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Erie Mfg., Co. |
| Location | Mchenry, IL, US |
| 510(k) history | 1 submissions · 1 cleared · 1978-1978 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k780359/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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