

K811397 ECAJul 2, 1981
45 days to decisionK811397 · Product code: **FOA** · General Hospital
Source: <https://www.510kdatabase.net/k811397/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Board, Cardiopulmonary (FOA) |
| Date received | May 18, 1981 |
| Decision date | Jul 2, 1981 |
| Days to decision | 45 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Gillco, Inc. |
| Location | Walker, MI, US |
| 510(k) history | 1 submissions · 1 cleared · 1981-1981 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k811397/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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