

K820939 C-101-PJun 22, 1982
77 days to decisionK820939 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k820939/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Implantable Pacemaker Pulse-generator (DXY) |
| Date received | Apr 6, 1982 |
| Decision date | Jun 22, 1982 |
| Days to decision | 77 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Coratomic, Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 15 submissions · 14 cleared · 1976-1986 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820939/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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