

**K925248 MODIFICATION OF INTRA-AORTIC BALLOON
CATHETERS**Sep 27, 1993
346 days to decisionK925248 · Product code: **DSP** · Cardiovascular
Source: <https://www.510kdatabase.net/k925248/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Balloon, Intra-aortic And Control (DSP) |
| Date received | Oct 16, 1992 |
| Decision date | Sep 27, 1993 |
| Days to decision | 346 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---|
| Company | Kontron Instruments, Inc. |
| Location | Mchenry, IL, US |
| Contact | WAYNE P GRIFFIN |
| 510(k) history | 57 submissions · 57 cleared · 1981-1993 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925248/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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