

**K855089 PERCUTANEOUS GUIDEWIRE PLACEMENT SYRINGE
50505**Feb 18, 1986
60 days to decisionK855089 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k855089/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Wire, Guide, Catheter (DQX) |
| Date received | Dec 20, 1985 |
| Decision date | Feb 18, 1986 |
| Days to decision | 60 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Dip, Inc. |
| Location | Mchenry, IL, US |
| Contact | RONALD A WILLIAMS |
| 510(k) history | 56 submissions · 56 cleared · 1979-1997 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k855089/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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