

K890959 GUIDEWIRESep 12, 1989
197 days to decisionK890959 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k890959/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Wire, Guide, Catheter (DQX) |
| Date received | Feb 27, 1989 |
| Decision date | Sep 12, 1989 |
| Days to decision | 197 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Guidewire Technologies, Inc. |
| Location | Salem, NH, US |
| Contact | DOUGLAS J CURTIS |
| 510(k) history | 3 submissions · 3 cleared · 1989-1990 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890959/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026