

K954772 QUICKELS QS 200 SYSTEMJul 11, 1996
272 days to decisionK954772 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k954772/>**SUBMISSION DETAILS**

| | |
|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received | Oct 13, 1995 |
| Decision date | Jul 11, 1996 |
| Days to decision | 272 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Ferguson Medical |
| Location | Chico, CA, US |
| Contact | FRANK FERGUSON |
| 510(k) history | 57 submissions · 55 cleared · 1985-2004 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k954772/> 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 May 18, 2026