

**K900882 MULTI DOPPLEX, MD-1**Aug 22, 1990  
177 days to decisionK900882 · Product code: **JAF** · Radiology  
Source: <https://www.510kdatabase.net/k900882/>**SUBMISSION DETAILS**

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|-----------------------|-------------------------------------|
| Decision              | Substantially Equivalent (Cleared)  |
| Submission type       | Traditional                         |
| Device classification | Monitor, Ultrasonic, Nonfetal (JAF) |
| Date received         | Feb 26, 1990                        |
| Decision date         | Aug 22, 1990                        |
| Days to decision      | 177 days                            |
| Third-party review    | No                                  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Huntleigh Technology, Inc.</b>       |
| Location       | Walker, MI, US                          |
| Contact        | JAMES BRITTON                           |
| 510(k) history | 23 submissions · 23 cleared · 1981-1999 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900882/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 20, 2026