

**K954127 BIOFEEDBACK DEVICE**Nov 14, 1995  
75 days to decisionK954127 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k954127/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Device, Biofeedback (HCC)          |
| Date received         | Aug 31, 1995                       |
| Decision date         | Nov 14, 1995                       |
| Days to decision      | 75 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>Verimed Holdings, Inc.</b>           |
| Location       | Mchenry, IL, US                         |
| Contact        | CHRISTOPHER J CHASE                     |
| 510(k) history | 12 submissions · 12 cleared · 1983-1998 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954127/> 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 June 30, 2026