

**K851732 CARBOCONE**Sep 16, 1985  
144 days to decisionK851732 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k851732/>**SUBMISSION DETAILS**

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|-----------------------|-------------------------------------|
| Decision              | Substantially Equivalent (Cleared)  |
| Submission type       | Traditional                         |
| Device classification | Electrode, Electrocardiograph (DRX) |
| Date received         | Apr 25, 1985                        |
| Decision date         | Sep 16, 1985                        |
| Days to decision      | 144 days                            |
| Third-party review    | No                                  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Imtech Corp.</b>                   |
| Location       | New York, NY, US                      |
| Contact        | PAUL NOBLE                            |
| 510(k) history | 1 submissions · 1 cleared · 1985-1985 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k851732/>; 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 July 8, 2026