

**K780179 LIQUID SCINTILLATION COUNTER**Feb 23, 1978  
20 days to decisionK780179 · Product code: **JJJ** · Toxicology  
Source: <https://www.510kdatabase.net/k780179/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)           |
| Submission type       | Traditional                                  |
| Device classification | Counter (beta, Gamma) For Clinical Use (JJJ) |
| Date received         | Feb 3, 1978                                  |
| Decision date         | Feb 23, 1978                                 |
| Days to decision      | 20 days                                      |
| Third-party review    | No   |

**APPLICANT**

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| Company        | <b>Roche Medical Electronics, Inc.</b>  |
| Location       | Mchenry, IL, US                         |
| 510(k) history | 16 submissions · 16 cleared · 1976-1979 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k780179/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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