

**K780341 SCINTILLATION, LIQUID, RACKBETA**May 26, 1978  
86 days to decisionK780341 · Product code: **JJJ** · Radiology  
Source: <https://www.510kdatabase.net/k780341/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter (beta, Gamma) For Clinical Use (JJJ)
Date received	Mar 1, 1978
Decision date	May 26, 1978
Days to decision	86 days
Third-party review	No

**APPLICANT**

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Company	<b>Lkb Instruments, Inc.</b>
Location	Mchenry, IL, US
510(k) history	52 submissions · 52 cleared · 1976-1987

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Device record: <https://www.510kdatabase.net/k780341/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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