FDA 510(k) Premarket Notification

K780341 SCINTILLATION, LIQUID, RACKBETA

CLEARED

May 26, 1978
86 days to decision

K780341 · Product code: **JJJ** · Radiology
Source: https://www.510kdatabase.net/k780341/

SUBMISSION DETAILS

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

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