

K830751 KEMTEK 102Apr 6, 1983
27 days to decisionK830751 · Product code: **JJJ** · Chemistry
Source: <https://www.510kdatabase.net/k830751/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter (beta, Gamma) For Clinical Use (JJJ)
Date received	Mar 10, 1983
Decision date	Apr 6, 1983
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Kemble Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1982-1984

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Device record: <https://www.510kdatabase.net/k830751/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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