FDA 510(k) Premarket Notification

K830751 KEMTEK 102

CLEARED

Apr 6, 1983
27 days to decision

K830751 · 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