FDA 510(k) Premarket Notification

K801259 NEEDLE COUNTER-SURE COUNT/100 COUNT

CLEARED

Jun 9, 1980
12 days to decision

K801259 · Product code: **MMK** · General Hospital
Source: https://www.510kdatabase.net/k801259/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Sharps (MMK)
Date received	May 28, 1980
Decision date	Jun 9, 1980
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Kleen Test Products**
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1976-1989