FDA 510(k) Premarket Notification

K820299 PROMPT INOCULATION SYSTEM

CLEARED

Mar 15, 1982
39 days to decision

K820299 · Product code: **LIE** · Microbiology
Source: https://www.510kdatabase.net/k820299/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reagent/device, Inoculum Calibration (LIE)
Date received	Feb 4, 1982
Decision date	Mar 15, 1982
Days to decision	39 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025