FDA 510(k) Premarket Notification

K911059 TURBIDOMETRIC STANDARD

CLEARED

Jun 10, 1991
91 days to decision

K911059 · Product code: **LIE** · Microbiology
Source: https://www.510kdatabase.net/k911059/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reagent/device, Inoculum Calibration (LIE)
Date received	Mar 11, 1991
Decision date	Jun 10, 1991
Days to decision	91 days
Third-party review	No

APPLICANT

Company	**Scientific Device Laboratory, Inc.**
Location	Mchenry, IL, US
Contact	STEWART LIPTON
510(k) history	28 submissions · 28 cleared · 1984-1994