FDA 510(k) Premarket Notification

K914170 LIQUICHEK THERAPEUTIC DRUG MONITORING CONTROL(TDM)

CLEARED

Nov 22, 1991
66 days to decision

K914170 · Product code: **DIF** · Toxicology
Source: https://www.510kdatabase.net/k914170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Sep 17, 1991
Decision date	Nov 22, 1991
Days to decision	66 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bio-Rad**
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510(k) history	319 submissions · 319 cleared · 1976-2017