FDA 510(k) Premarket Notification

K992033 MODIFICATION TO 'RAPID DRUG SCREEN' 9-PANEL

CLEARED

Jun 30, 1999
14 days to decision

K992033 · Product code: **DKZ** · Toxicology
Source: https://www.510kdatabase.net/k992033/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Amphetamine (DKZ)
Date received	Jun 16, 1999
Decision date	Jun 30, 1999
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**American Bio Medica Corp.**
Location	Washington, DC, US
Contact	JOHN B DUBECK
510(k) history	30 submissions · 30 cleared · 1997-2017