

K993961 RAPID DRUG SCREEN 3-PANEL TEST FOR COCAINE, MARIJUANA (THC) (CANNABINOIDS) AND OPIATES (3 PANEL TEST FOR CTO)

Jan 24, 2000
63 days to decision

K993961 · Product code: LDJ · Toxicology
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SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Cannabinoids (LDJ)
Date received	Nov 22, 1999
Decision date	Jan 24, 2000
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k993961/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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