

K981705 'RAPID DRUG SCREEN'; 8 PANEL-LOW VOLUME

Aug 27, 1998
105 days to decision

K981705 · Product code: **DKZ** · Toxicology
Source: <https://www.510kdatabase.net/k981705/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Amphetamine (DKZ)
Date received	May 14, 1998
Decision date	Aug 27, 1998
Days to decision	105 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k981705/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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