

**K030835 &apos;RAPID ONE&apos; - PROPOXYPHENE TEST**May 22, 2003  
66 days to decisionK030835 · Product code: **JXN** · Toxicology  
Source: <https://www.510kdatabase.net/k030835/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Propoxyphene (JXN)
Date received	Mar 17, 2003
Decision date	May 22, 2003
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030835/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 9, 2026