

**K040029 SAFETYNET GUIDEWIRE INTRODUCER**Apr 5, 2004  
88 days to decisionK040029 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k040029/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jan 8, 2004
Decision date	Apr 5, 2004
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Injectimed, Inc.</b>
Location	Ventura, CA, US
Contact	GEORGE C BRDLIK
510(k) history	3 submissions · 3 cleared · 1992-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k040029/> 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 25, 2026