

**K052022 ASAHI PTCA GUIDE WIRE FIELDER**Aug 25, 2005  
30 days to decisionK052022 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k052022/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 26, 2005
Decision date	Aug 25, 2005
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Asahi Intecc Co., Ltd.</b>
Location	Seto-Shi, JP
Contact	YOSHI TERAJ
Website	<a href="https://www.asahi-intecc.com">https://www.asahi-intecc.com</a>
510(k) history	84 submissions · 84 cleared · 2003-2026

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052022/> 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 28, 2026