

K013024 SELECTIVA GUIDEWIREDec 4, 2001
88 days to decisionK013024 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k013024/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Sep 7, 2001
Decision date	Dec 4, 2001
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Neometrics, Inc.
Location	Mchenry, IL, US
Contact	GENE CHAMPEAU
510(k) history	21 submissions · 21 cleared · 1978-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013024/> Data sourced from FDA 510(k) public records (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. - Generated June 3, 2026