FDA 510(k) Premarket Notification

K013024 SELECTIVA GUIDEWIRE

CLEARED

Dec 4, 2001
88 days to decision

K013024 · 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