

K920408 DESERET GUIDEWIRE INTRODUCER IIIApr 7, 1992
67 days to decisionK920408 · Product code: **DQO** · Cardiovascular
Source: <https://www.510kdatabase.net/k920408/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 31, 1992
Decision date	Apr 7, 1992
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bd Becton Dickinson Vacutainer Systems Preanalytic
Location	Washington, DC, US
Contact	CHARLES J WELLE
510(k) history	632 submissions · 625 cleared · 1976-2001

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