

**K020012 MODIFICATION TO RENEGADE FIBER BRAIDED
MICROCATHETER**Jan 29, 2002
26 days to decisionK020012 · Product code: **DQO** · Cardiovascular
Source: <https://www.510kdatabase.net/k020012/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 3, 2002
Decision date	Jan 29, 2002
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Boston Scientific/Medi-Tech
Location	Natick,, MA, US
Contact	JODI L GREENIZEN
510(k) history	5 submissions · 5 cleared · 2001-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020012/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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