

K252390 Telescope Guide Extension CatheterOct 29, 2025
90 days to decisionK252390 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k252390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jul 31, 2025
Decision date	Oct 29, 2025
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic, Ireland
Location	Shoreview, MN, US
Contact	Denise Keehan
510(k) history	7 submissions · 7 cleared · 2006-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k252390/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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