

K172995 Destination Carotid Guiding Sheath, Destination Peripheral Guiding Sheath, Destination Renal Guiding SheathNov 21, 2017
55 days to decisionK172995 · Product code: **DYB** · Cardiovascular
Source: <https://www.510kdatabase.net/k172995/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Sep 27, 2017
Decision date	Nov 21, 2017
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Terumo Medical Corporation
Location	Elkton, MD, US
Contact	Liang Lu
510(k) history	14 submissions · 14 cleared · 2011-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k172995/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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