

K820859 COAXIAL DUAL FLOW CATHETERApr 16, 1982
18 days to decisionK820859 · Product code: **LBW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k820859/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Dialysis, Single Needle (co-axial Flow) (LBW)
Date received	Mar 29, 1982
Decision date	Apr 16, 1982
Days to decision	18 days
Third-party review	No

APPLICANT

Company	Terumo Medical Corp.
Location	Elkton, MD, US
510(k) history	143 submissions · 143 cleared · 1980-2011

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Device record: <https://www.510kdatabase.net/k820859/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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