

K834352 ADULT VENT CATHETERJan 27, 1984
45 days to decisionK834352 · Product code: **DRA** · CardiovascularSource: <https://www.510kdatabase.net/k834352/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Steerable (DRA)
Date received	Dec 13, 1983
Decision date	Jan 27, 1984
Days to decision	45 days
Third-party review	No

APPLICANT

Company	Dlp, Inc.
Location	Mchenry, IL, US
510(k) history	56 submissions · 56 cleared · 1979-1997

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Device record: <https://www.510kdatabase.net/k834352/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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