

K832023 EMBOLECTOMY CATHETERAug 31, 1983
79 days to decisionK832023 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k832023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Jun 13, 1983
Decision date	Aug 31, 1983
Days to decision	79 days
Third-party review	No

APPLICANT

Company	Bionomy, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1981-1985

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Device record: <https://www.510kdatabase.net/k832023/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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