

K801173 ANGIO-FLO MONITORING KITMay 28, 1980
12 days to decisionK801173 · Product code: **KRA** · CardiovascularSource: <https://www.510kdatabase.net/k801173/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	May 16, 1980
Decision date	May 28, 1980
Days to decision	12 days
Third-party review	No

APPLICANT

Company	Hospital Products Co.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1983

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Device record: <https://www.510kdatabase.net/k801173/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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