

K800726 EYEWI ANKORSMay 8, 1980
37 days to decisionK800726 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k800726/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Apr 1, 1980
Decision date	May 8, 1980
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Eyevi & Co.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800726/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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