FDA 510(k) Premarket Notification

K800726 EYEWI ANKORS

CLEARED

May 8, 1980
37 days to decision

K800726 · 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