

K842999 VENI-SHIELDSep 24, 1984
56 days to decisionK842999 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k842999/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Jul 30, 1984
Decision date	Sep 24, 1984
Days to decision	56 days
Third-party review	No

APPLICANT

Company	U.S. Mfg. & Management, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 3 cleared · 1984-1992

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Device record: <https://www.510kdatabase.net/k842999/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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