

K832300 I.V. FILTER HOLDEROct 4, 1983
83 days to decisionK832300 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k832300/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Jul 13, 1983
Decision date	Oct 4, 1983
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Saxnor Corp.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k832300/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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