

K812080 AEGIS I.V. STRIPSAug 3, 1981
12 days to decisionK812080 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k812080/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Device, Intravascular Catheter Securement (KMK) |
| Date received | Jul 22, 1981 |
| Decision date | Aug 3, 1981 |
| Days to decision | 12 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Warner-Lambert Co. |
| Location | Mchenry, IL, US |
| 510(k) history | 50 submissions · 50 cleared · 1979-2003 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812080/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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