

K890379 UTERINE TENTACULUMFeb 13, 1989
21 days to decisionK890379 · Product code: **HDC** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k890379/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Tenaculum, Uterine (HDC) |
| Date received | Jan 23, 1989 |
| Decision date | Feb 13, 1989 |
| Days to decision | 21 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Kinetic Medical Products |
| Location | Erie, PA, US |
| Contact | JAMES I LAUGHNER |
| 510(k) history | 63 submissions · 63 cleared · 1989-1989 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890379/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026