

K841682 UTERINE INJECTOR 4.0MMJul 3, 1984
71 days to decisionK841682 · Product code: **HES** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k841682/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Insufflator, Carbon-dioxide, Uterotubal (and Accessories) (HES) |
| Date received | Apr 23, 1984 |
| Decision date | Jul 3, 1984 |
| Days to decision | 71 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Humi Corp. |
| Location | Walker, MI, US |
| 510(k) history | 2 submissions · 1 cleared · 1984-1984 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k841682/> 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 June 30, 2026