

K122820 VIKY UPJun 18, 2013
277 days to decisionK122820 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k122820/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Cannula, Manipulator/injector, Uterine (LKF) |
| Date received | Sep 14, 2012 |
| Decision date | Jun 18, 2013 |
| Days to decision | 277 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Endocontrol |
| Location | La Tronche, FR |
| Contact | HUGUEL CARINE |
| 510(k) history | 2 submissions · 2 cleared · 2008-2013 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122820/> 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 8, 2026