

K102815 SURELIFT PROLAPSE SYSTEMJul 7, 2011
282 days to decisionK102815 · Product code: **OTP** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k102815/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Mesh, Surgical, Synthetic, Urogynecologic, For Pelvic Organ Prolapse, Transvaginally Placed (OTP) |
| Date received | Sep 28, 2010 |
| Decision date | Jul 7, 2011 |
| Days to decision | 282 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Neomedic International |
| Location | Minneapolis, MN, US |
| Contact | JEFFREY R SHIDEMAN |
| 510(k) history | 1 submissions · 1 cleared · 2011-2011 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102815/> 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 19, 2026