

K042341 ULTRACLIP II USSep 20, 2004
21 days to decisionK042341 · Product code: **NEU** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k042341/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Marker, Radiographic, Implantable (NEU) |
| Date received | Aug 30, 2004 |
| Decision date | Sep 20, 2004 |
| Days to decision | 21 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---|
| Company | Inrad |
| Location | Kentwood, MI, US |
| Contact | MELISSA LALOMIA |
| 510(k) history | 11 submissions · 11 cleared · 1998-2009 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042341/> 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 22, 2026