

K052830 SPORTMESHJan 19, 2006
106 days to decisionK052830 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k052830/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Mesh, Surgical, Polymeric (FTL) |
| Date received | Oct 5, 2005 |
| Decision date | Jan 19, 2006 |
| Days to decision | 106 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Artimplant AB |
| Location | Washington, DC, US |
| Contact | TERRY SHERIDAN POWELL |
| 510(k) history | 7 submissions · 7 cleared · 2003-2007 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052830/> 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 19, 2026