

K874383 MENTOR PLUS (CONDOM)Mar 7, 1988
133 days to decisionK874383 · Product code: **HIS** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k874383/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Condom (HIS) |
| Date received | Oct 26, 1987 |
| Decision date | Mar 7, 1988 |
| Days to decision | 133 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Arcon Corp. |
| Location | Stewartville, MN, US |
| Contact | RICHARD FRYAR |
| 510(k) history | 2 submissions · 2 cleared · 1984-1988 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k874383/>; 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