

K820123 SUNFLOWERMar 1, 1982
42 days to decisionK820123 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820123/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Light, Ultraviolet, Dermatological (FTC) |
| Date received | Jan 18, 1982 |
| Decision date | Mar 1, 1982 |
| Days to decision | 42 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Solana, Inc. |
| Location | Walker, MI, US |
| 510(k) history | 1 submissions · 1 cleared · 1982-1982 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k820123/> 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 July 9, 2026