

**K860347 SUNLAMP MODELS 24N, 30R, 30RE, 37RE**Feb 12, 1986  
14 days to decisionK860347 · Product code: **FTC** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k860347/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Light, Ultraviolet, Dermatological (FTC) |
| Date received         | Jan 29, 1986                             |
| Decision date         | Feb 12, 1986                             |
| Days to decision      | 14 days                                  |
| Third-party review    | No                                       |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Gti, Inc.</b>                      |
| Location       | East Lyme, CT, US                     |
| Contact        | GERHARD FRANZ                         |
| 510(k) history | 2 submissions · 2 cleared · 1986-1992 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860347/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 4, 2026