FDA 510(k) Premarket Notification

# K883563 SHARPLAN 2100 ND:YAG LASER, SERIES 2900 SYN. TIPS

**CLEARED**

Sep 27, 1988
39 days to decision

K883563 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k883563/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Powered Laser Surgical Instrument (GEX) |
| Date received | Aug 19, 1988 |
| Decision date | Sep 27, 1988 |
| Days to decision | 39 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Sharplan Lasers, Inc.** |
| Location | Allendale, NJ, US |
| Contact | STEPHEN DALTON |
| 510(k) history | 78 submissions · 78 cleared · 1986-1997 |