FDA 510(k) Premarket Notification

# K073503 VERSA LITE

**CLEARED**

Jan 23, 2008
41 days to decision

K073503 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k073503/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Light, Ultraviolet, Dermatological (FTC) |
| Date received | Dec 13, 2007 |
| Decision date | Jan 23, 2008 |
| Days to decision | 41 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Diomedics, Inc.** |
| Location | The Woodlands, TX, US |
| Contact | GARY MOCNIK |
| 510(k) history | 3 submissions · 3 cleared · 1993-2008 |