FDA 510(k) Premarket Notification

# K893949 #157 EXAM LIGHT

**CLEARED**

Aug 7, 1989
67 days to decision

K893949 · Product code: **KFK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k893949/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Saw, Pneumatically Powered (KFK) |
| Date received | Jun 1, 1989 |
| Decision date | Aug 7, 1989 |
| Days to decision | 67 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Medmark, Inc.** |
| Location | Versailles, OH, US |
| Contact | JOHN OLDIGES |
| 510(k) history | 3 submissions · 3 cleared · 1989-1991 |