

K840371 FLEXIBLE ILLUMINATOR 5Apr 24, 1984
88 days to decisionK840371 · Product code: **FTF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k840371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Illuminator, Non-remote (FTF)
Date received	Jan 27, 1984
Decision date	Apr 24, 1984
Days to decision	88 days
Third-party review	No

APPLICANT

Company	Suncoast Medical Manufacturers, Inc.
Location	Mchenry, IL, US
510(k) history	31 submissions · 29 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840371/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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