FDA 510(k) Premarket Notification

K840371 FLEXIBLE ILLUMINATOR 5

CLEARED

Apr 24, 1984
88 days to decision

K840371 · Product code: **FTF** · General & Plastic Surgery
Source: 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