

K820492 SPECTRA305Mar 31, 1982
36 days to decisionK820492 · Product code: **FTC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820492/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 23, 1982
Decision date	Mar 31, 1982
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Ultralite
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1977-1991

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Device record: <https://www.510kdatabase.net/k820492/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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