

K827890 FOLDALITE UNITAug 23, 1982
12 days to decisionK827890 · Product code: **FTC** · Radiology
Source: <https://www.510kdatabase.net/k827890/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Aug 11, 1982
Decision date	Aug 23, 1982
Days to decision	12 days
Third-party review	No

APPLICANT

Company	National Biological Corp.
Location	Walker, MI, US
510(k) history	27 submissions · 27 cleared · 1977-2007

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Device record: <https://www.510kdatabase.net/k827890/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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