FDA 510(k) Premarket Notification

K864810 LIDSPLINT

CLEARED

Jan 20, 1987
42 days to decision

K864810 · Product code: **HMP** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k864810/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pad, Eye (HMP)
Date received	Dec 9, 1986
Decision date	Jan 20, 1987
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Precision Therapeutics, Inc.**
Location	Las Vegas, NV, US
Contact	LEONARDI, M.D.
510(k) history	2 submissions · 2 cleared · 1987-1987