

K864810 LIDSPLINTJan 20, 1987
42 days to decisionK864810 · Product code: **HMP** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k864810/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026