

**K861746 KEELER DUAL LIGHTSOURCE**Aug 21, 1986  
107 days to decisionK861746 · Product code: **HPQ** · Ophthalmic  
Source: <https://www.510kdatabase.net/k861746/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Headlamp, Operating, Ac-powered (HPQ)
Date received	May 6, 1986
Decision date	Aug 21, 1986
Days to decision	107 days
Third-party review	No

**APPLICANT**

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Company	<b>Keeler Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861746/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 20, 2026