

**K860062 OCULAB PACHPEN**Mar 10, 1986  
63 days to decisionK860062 · Product code: **IYO** · Radiology  
Source: <https://www.510kdatabase.net/k860062/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jan 6, 1986
Decision date	Mar 10, 1986
Days to decision	63 days
Third-party review	No

**APPLICANT**

---

Company	<b>Oculab, Inc.</b>
Location	Gendale, CA, US
Contact	WALLACE, M.D.
510(k) history	7 submissions · 7 cleared · 1985-1989

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860062/>; 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 May 21, 2026