FDA 510(k) Premarket Notification

K860062 OCULAB PACHPEN

CLEARED

Mar 10, 1986
63 days to decision

K860062 · 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	**Oculab, Inc.**
Location	Gendale, CA, US
Contact	WALLACE, M.D.
510(k) history	7 submissions · 7 cleared · 1985-1989