FDA 510(k) Premarket Notification

K874555 KEELER ACU 22XT OPHTHALMIC CRYO UNIT

CLEARED

Jan 29, 1988
86 days to decision

K874555 · Product code: **HPS** · Ophthalmic
Source: https://www.510kdatabase.net/k874555/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryophthalmic (HPS)
Date received	Nov 4, 1987
Decision date	Jan 29, 1988
Days to decision	86 days
Third-party review	No

APPLICANT

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