FDA 510(k) Premarket Notification

K062412 CRYOMATIC

CLEARED

Nov 6, 2006
81 days to decision

K062412 · Product code: **HRN** · Ophthalmic
Source: https://www.510kdatabase.net/k062412/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryophthalmic, Ac-powered (HRN)
Date received	Aug 17, 2006
Decision date	Nov 6, 2006
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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