

K062412 CRYOMATICNov 6, 2006
81 days to decisionK062412 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062412/> Data sourced from FDA 510(k) public records (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. - Generated June 20, 2026