

K830366 DIAMOND KNIFEMar 9, 1983
33 days to decisionK830366 · Product code: **HNN** · Ophthalmic
Source: <https://www.510kdatabase.net/k830366/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Knife, Ophthalmic (HNN)
Date received	Feb 4, 1983
Decision date	Mar 9, 1983
Days to decision	33 days
Third-party review	No

APPLICANT

Company	Koi, Inc.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1981-1986

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k830366/> 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 May 21, 2026