

**K812450 KOI DIAMOND KNIFE**Sep 21, 1981  
24 days to decisionK812450 · Product code: **HNN** · Ophthalmic  
Source: <https://www.510kdatabase.net/k812450/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Knife, Ophthalmic (HNN)
Date received	Aug 28, 1981
Decision date	Sep 21, 1981
Days to decision	24 days
Third-party review	No

**APPLICANT**

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

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Device record: <https://www.510kdatabase.net/k812450/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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