FDA 510(k) Premarket Notification

K812450 KOI DIAMOND KNIFE

CLEARED

Sep 21, 1981
24 days to decision

K812450 · Product code: **HNN** · Ophthalmic
Source: https://www.510kdatabase.net/k812450/

SUBMISSION DETAILS

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

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