

K812010 KOI LACRIMAL INTUBATION SYSTEMJul 28, 1981
11 days to decisionK812010 · Product code: **HMX** · Ophthalmic
Source: <https://www.510kdatabase.net/k812010/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	Jul 17, 1981
Decision date	Jul 28, 1981
Days to decision	11 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k812010/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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