

K813314 NLD IMPLANT SETFeb 24, 1982
92 days to decisionK813314 · Product code: **HMX** · Ophthalmic
Source: <https://www.510kdatabase.net/k813314/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	Nov 24, 1981
Decision date	Feb 24, 1982
Days to decision	92 days
Third-party review	No

APPLICANT

Company	Design Research Assoc., Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1982

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Device record: <https://www.510kdatabase.net/k813314/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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