FDA 510(k) Premarket Notification

K813314 NLD IMPLANT SET

CLEARED

Feb 24, 1982
92 days to decision

K813314 · 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