FDA 510(k) Premarket Notification

K843039 SITE ANTERIOR CAPSULOTOMER

CLEARED

Nov 21, 1984
111 days to decision

K843039 · Product code: **HMX** · Ophthalmic
Source: https://www.510kdatabase.net/k843039/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	Aug 2, 1984
Decision date	Nov 21, 1984
Days to decision	111 days
Third-party review	No

APPLICANT

Company	**Chiron Vision Corp.**
Location	Irvine, CA, US
Contact	TICKNOR
510(k) history	34 submissions · 34 cleared · 1980-1997