FDA 510(k) Premarket Notification

K840063 CATAREX VITRECUTTER I/A DISPOS. CUT

CLEARED

Oct 5, 1984
270 days to decision

K840063 · Product code: **HQC** · Ophthalmic
Source: https://www.510kdatabase.net/k840063/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Jan 9, 1984
Decision date	Oct 5, 1984
Days to decision	270 days
Third-party review	No

APPLICANT

Company	**Calliope Corp.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1983-1984