FDA 510(k) Premarket Notification

K840660 POSTERIOR CAPSULOTOMY CYSTITOME

CLEARED

Mar 16, 1984
30 days to decision

K840660 · Product code: **HNY** · Ophthalmic
Source: https://www.510kdatabase.net/k840660/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystotome (HNY)
Date received	Feb 15, 1984
Decision date	Mar 16, 1984
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Sharpoint, Inc.**
Location	Walker, MI, US
510(k) history	9 submissions · 9 cleared · 1982-1986