FDA 510(k) Premarket Notification

K802593 KERATOPLASTY SUTURING LENS

CLEARED

Dec 22, 1980
62 days to decision

K802593 · Product code: **HPX** · Ophthalmic
Source: https://www.510kdatabase.net/k802593/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Contact (polymethylmethacrylate) (HPX)
Date received	Oct 21, 1980
Decision date	Dec 22, 1980
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Paris Contact Lens Laboratory**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k802593/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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