FDA 510(k) Premarket Notification

K900080 NEW IOWA MOTILITY IMPLANT

CLEARED

Mar 23, 1990
78 days to decision

K900080 · Product code: **HPZ** · Ophthalmic
Source: https://www.510kdatabase.net/k900080/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Eye Sphere (HPZ)
Date received	Jan 4, 1990
Decision date	Mar 23, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Oculo Plastik, Inc.**
Location	Montreal, Quebec Canada, CA
Contact	JEAN-FRANCOIS DURETT
510(k) history	13 submissions · 13 cleared · 1990-2013