FDA 510(k) Premarket Notification

K900927 CODERE-DURETTE ORBITAL FLOOR IMPLANT

CLEARED

May 15, 1990
77 days to decision

K900927 · Product code: **HQX** · Ophthalmic
Source: https://www.510kdatabase.net/k900927/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Orbital, Extra-ocular (HQX)
Date received	Feb 27, 1990
Decision date	May 15, 1990
Days to decision	77 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