

**K900927 CODERE-DURETTE ORBITAL FLOOR IMPLANT**May 15, 1990  
77 days to decisionK900927 · Product code: **HQX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k900927/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900927/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 9, 2026