

**K821381 IOL FORCEPS**Jun 14, 1982  
35 days to decisionK821381 · Product code: **KYB** · Ophthalmic  
Source: <https://www.510kdatabase.net/k821381/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	May 10, 1982
Decision date	Jun 14, 1982
Days to decision	35 days
Third-party review	No

**APPLICANT**

---

Company	<b>Cilco, Inc.</b>
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1982-1986

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k821381/> 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 May 21, 2026