

K820933 CILCO I/A HANDPIECEJun 14, 1982
73 days to decisionK820933 · Product code: **HQE** · Ophthalmic
Source: <https://www.510kdatabase.net/k820933/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Apr 2, 1982
Decision date	Jun 14, 1982
Days to decision	73 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k820933/> Data sourced from FDA 510(k) public records (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. - Generated May 21, 2026