FDA 510(k) Premarket Notification

K820933 CILCO I/A HANDPIECE

CLEARED

Jun 14, 1982
73 days to decision

K820933 · 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