

K801850 CHARLES PNEUMATIC INTRAOCULAR SCISSORSSep 16, 1980
43 days to decisionK801850 · Product code: **HQE** · Ophthalmic
Source: <https://www.510kdatabase.net/k801850/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Aug 4, 1980
Decision date	Sep 16, 1980
Days to decision	43 days
Third-party review	No

APPLICANT

Company	Cooper Medical Devices Corp.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1979-1981

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801850/> 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 June 30, 2026