

**K840063 CATAREX VITRECUITTER I/A DISPOS. CUT**Oct 5, 1984  
270 days to decisionK840063 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k840063/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Jan 9, 1984
Decision date	Oct 5, 1984
Days to decision	270 days
Third-party review	No

**APPLICANT**

---

Company	<b>Calliope Corp.</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1983-1984

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k840063/> 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 July 1, 2026