

**K031735 CENTURION SES EPIKERATOME**Aug 21, 2003  
78 days to decisionK031735 · Product code: **HNO** · Ophthalmic  
Source: <https://www.510kdatabase.net/k031735/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Ac-powered (HNO)
Date received	Jun 4, 2003
Decision date	Aug 21, 2003
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ciba Vision Corporation</b>
Location	Duluth, GA, US
Contact	PENNY NORTHCUTT
510(k) history	44 submissions · 44 cleared · 1994-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031735/> 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 June 7, 2026