

**K010534 CROSS-CHECKS DUAL, MODEL CI 125**Aug 6, 2001  
164 days to decisionK010534 · Product code: **JOJ** · General Hospital  
Source: <https://www.510kdatabase.net/k010534/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Physical/chemical Sterilization Process (JOJ)
Date received	Feb 23, 2001
Decision date	Aug 6, 2001
Days to decision	164 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Steritec Products, Inc.</b>
Location	Castle Rock, CO, US
Contact	LINDA NELSON
510(k) history	12 submissions · 12 cleared · 2000-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010534/> 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 23, 2026