

**K020470 MODIFICATION TO CORLINK AUTOMATED ANASTOMOTIC DEVICE**Mar 8, 2002  
24 days to decision

K020470 · Product code: FZP · General &amp; Plastic Surgery

Source: <https://www.510kdatabase.net/k020470/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Clip, Implantable (FZP)
Date received	Feb 12, 2002
Decision date	Mar 8, 2002
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bypass , Ltd.</b>
Location	Washington, DC, US
Contact	JONATHAN S KAHAN
510(k) history	2 submissions · 2 cleared · 2001-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020470/>. 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 3, 2026