

**K041370 LIFESTENT LP SDS BILIARY ENDOPROSTHESIS**Jun 24, 2004  
31 days to decisionK041370 · Product code: **FGE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k041370/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Special
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	May 24, 2004
Decision date	Jun 24, 2004
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Orbus Medical Technologies</b>
Location	Fort Lauderdale, FL, US
Contact	JIM CLOSSICK
510(k) history	8 submissions · 0 cleared · 2001-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041370/> 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 30, 2026