

**K072050 PROLONG-EX, MODELS: PLEX-50CS, PLEX-100CS,  
PLEX-150CS**Oct 18, 2007  
84 days to decisionK072050 · Product code: **BSO** · Anesthesiology  
Source: <https://www.510kdatabase.net/k072050/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Conduction, Anesthetic (BSO)
Date received	Jul 26, 2007
Decision date	Oct 18, 2007
Days to decision	84 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Life-Tech, Inc.</b>
Location	Stafford, TX, US
Contact	JEFF KASOFF
510(k) history	14 submissions · 14 cleared · 2005-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072050/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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