

**K021480 EXTENSION SET LIGHT-SAFE, MODEL BC565**Jun 27, 2002  
50 days to decisionK021480 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k021480/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 8, 2002
Decision date	Jun 27, 2002
Days to decision	50 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Codan US Corp.</b>
Location	Irvine, CA, US
Contact	MARILYN R POURAZAR
510(k) history	7 submissions · 7 cleared · 1999-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021480/> 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 15, 2026