

**K152157 LC Bead LUMI**Dec 11, 2015  
130 days to decisionK152157 · Product code: **KRD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k152157/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Aug 3, 2015
Decision date	Dec 11, 2015
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biocompatibles UK , Ltd.</b>
Location	Farnham, GB
Contact	April Kent
510(k) history	2 submissions · 2 cleared · 2015-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k152157/> 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 3, 2026