

**K072114 IMPULSE 6000D/7000DP**Jan 28, 2008  
180 days to decisionK072114 · Product code: **DRL** · Cardiovascular  
Source: <https://www.510kdatabase.net/k072114/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Defibrillator (DRL)
Date received	Aug 1, 2007
Decision date	Jan 28, 2008
Days to decision	180 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fluke Biomedical</b>
Location	Orange, CA, US
Contact	COLLEEN BOSWELL
510(k) history	6 submissions · 6 cleared · 2008-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072114/> 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 9, 2026