

**K093813 OCCLUSIN 500 ARTIFICIAL EMBOLIZATION DEVICE**May 10, 2010  
150 days to decisionK093813 · Product code: **KRD** · CardiovascularSource: <https://www.510kdatabase.net/k093813/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Dec 11, 2009
Decision date	May 10, 2010
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Imbiotechnologies</b>
Location	North Attleboro, MA, US
Contact	ROSINA ROBINSON
510(k) history	1 submissions · 1 cleared · 2010-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093813/> 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 21, 2026