

**K060929 IMMULITE 2500 VITAMIN B12 MODEL L5KVB2-200  
TEST,L5KVB6-600 TEST, IMMULITE 2500 FOLIC ACID MODEL  
L5KFO2-200 TEST,L5KFO6-6**

Apr 28, 2006  
23 days to decision

K060929 · Product code: **CDD** · Chemistry  
Source: <https://www.510kdatabase.net/k060929/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Radioassay, Vitamin B12 (CDD)
Date received	Apr 5, 2006
Decision date	Apr 28, 2006
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Diagnostic Products Corporation</b>
Location	Los Angeles, CA, US
Contact	DEBORAH L MORRIS
510(k) history	3 submissions · 3 cleared · 2006-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k060929/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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