

**K093815 LIGHT DIAGNOSTICS HUMAN METAPNEUMOVIRUS  
DFA KIT, MODEL CATALOG NUMBER 3124**Mar 12, 2010  
88 days to decisionK093815 · Product code: **OMG** · Microbiology  
Source: <https://www.510kdatabase.net/k093815/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, Human Metapneumovirus (OMG)
Date received	Dec 14, 2009
Decision date	Mar 12, 2010
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Millipore Corporation</b>
Location	Temecula, CA, US
Contact	CATHERINE KARAMAN
510(k) history	2 submissions · 2 cleared · 2009-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093815/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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