

**K822347 LIDOCAINE FLUORESCENT IMMUNOASSAY**Aug 25, 1982  
19 days to decisionK822347 · Product code: **KLR** · Toxicology  
Source: <https://www.510kdatabase.net/k822347/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Lidocaine (KLR)
Date received	Aug 6, 1982
Decision date	Aug 25, 1982
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>American Diagnostic Corp.</b>
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1980-2017

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

Device record: <https://www.510kdatabase.net/k822347/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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