

**K782159 DIAGNOSTIC KIT, FERRITAB-FE-59**Feb 1, 1979  
41 days to decisionK782159 · Product code: **JJA** · Chemistry  
Source: <https://www.510kdatabase.net/k782159/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radio-labeled Iron Method, Iron (non-heme) (JJA)
Date received	Dec 22, 1978
Decision date	Feb 1, 1979
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Leeco Diagnostics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	49 submissions · 49 cleared · 1979-1989

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Device record: <https://www.510kdatabase.net/k782159/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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