

**K830564 FEB TEST**Mar 29, 1983  
34 days to decisionK830564 · Product code: **CFM** · Chemistry  
Source: <https://www.510kdatabase.net/k830564/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bathophenanthroline, Colorimetry, Iron (non-heme) (CFM)
Date received	Feb 23, 1983
Decision date	Mar 29, 1983
Days to decision	34 days
Third-party review	No

**APPLICANT**

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Company	<b>Wako Chemicals USA, Inc.</b>
Location	Mchenry, IL, US
510(k) history	124 submissions · 123 cleared · 1981-2011

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

Device record: <https://www.510kdatabase.net/k830564/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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