

**K833854 LH RIA**Jan 13, 1984  
70 days to decisionK833854 · Product code: **CEP** · Chemistry  
Source: <https://www.510kdatabase.net/k833854/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Luteinizing Hormone (CEP)
Date received	Nov 4, 1983
Decision date	Jan 13, 1984
Days to decision	70 days
Third-party review	No

**APPLICANT**

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Company	<b>Syncor Intl. Corp.</b>
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1983-1995

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Device record: <https://www.510kdatabase.net/k833854/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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