

K840174 LH QUIK RIA KITMar 19, 1984
62 days to decisionK840174 · Product code: **CEP** · Chemistry
Source: <https://www.510kdatabase.net/k840174/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Luteinizing Hormone (CEP)
Date received	Jan 17, 1984
Decision date	Mar 19, 1984
Days to decision	62 days
Third-party review	No

APPLICANT

Company	Pacific Biotech, Inc.
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1983-1993

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Device record: <https://www.510kdatabase.net/k840174/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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